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Pan-European rural monitoring network shows dominance of NH_3 gas and NH_4NO_3 aerosol in inorganic atmospheric pollution load

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Abstract. A comprehensive European dataset on monthly atmospheric NH_3 , acid gases (HNO_3 , SO_2 , HCl), and aerosols (NH_4^+ , NO_3^- , SO_4^{2-} , Cl^- , Na^+ , Ca^{2+} , Mg^{2+}) is presented and analysed. Speciated measurements were made with a low-volume denuder and filter pack method (DENuder for Long-Term Atmospheric sampling, DELTA[®]) as part of the EU NitroEurope (NEU) integrated project. Altogether, there were 64 sites in 20 countries (2006–2010), coordinated between seven European laboratories. Bulk wet-deposition measurements were carried out at 16 co-located sites (2008–2010). Inter-comparisons of chemical analysis and DELTA[®] measurements allowed an assessment of comparability between laboratories.

The form and concentrations of the different gas and aerosol components measured varied between individual sites and grouped sites according to country, European re-

gions, and four main ecosystem types (crops, grassland, forests, and semi-natural). The smallest concentrations (with the exception of SO_4^{2-} and Na^+) were in northern Europe (Scandinavia), with broad elevations of all components across other regions. SO_2 concentrations were highest in central and eastern Europe, with larger SO_2 emissions, but particulate SO_4^{2-} concentrations were more homogeneous between regions. Gas-phase NH_3 was the most abundant single measured component at the majority of sites, with the largest variability in concentrations across the network. The largest concentrations of NH_3 , NH_4^+ , and NO_3^- were at cropland sites in intensively managed agricultural areas (e.g. Borgo Cioffi in Italy), and the smallest were at remote semi-natural and forest sites (e.g. Lompolojänkka, Finland), highlighting the potential for NH_3 to drive the formation of both NH_4^+ and NO_3^- aerosol. In the aerosol phase, NH_4^+ was highly corre-

lated with both NO₃⁻ and SO₄²⁻, with a near-1 : 1 relationship between the equivalent concentrations of NH₄⁺ and sum (NO₃⁻ + SO₄²⁻), of which around 60 % was as NH₄NO₃.

Distinct seasonality was also observed in the data, influenced by changes in emissions, chemical interactions, and the influence of meteorology on partitioning between the main inorganic gases and aerosol species. Springtime maxima in NH₃ were attributed to the main period of manure spreading, while the peak in summer and trough in winter were linked to the influence of temperature and rainfall on emissions, deposition, and gas–aerosol-phase equilibrium. Seasonality in SO₂ was mainly driven by emissions (combustion), with concentrations peaking in winter, except in southern Europe, where the peak occurred in summer. Particulate SO₄²⁻ showed large peaks in concentrations in summer in southern and eastern Europe, contrasting with much smaller peaks occurring in early spring in other regions. The peaks in particulate SO₄²⁻ coincided with peaks in NH₃ concentrations, attributed to the formation of the stable (NH₄)₂SO₄. HNO₃ concentrations were more complex, related to traffic and industrial emissions, photochemistry, and HNO₃:NH₄NO₃ partitioning. While HNO₃ concentrations were seen to peak in the summer in eastern and southern Europe (increased photochemistry), the absence of a spring peak in HNO₃ in all regions may be explained by the depletion of HNO₃ through reaction with surplus NH₃ to form the semi-volatile aerosol NH₄NO₃. Cooler, wetter conditions in early spring favour the formation and persistence of NH₄NO₃ in the aerosol phase, consistent with the higher springtime concentrations of NH₄⁺ and NO₃⁻. The seasonal profile of NO₃⁻ was mirrored by NH₄⁺, illustrating the influence of gas–aerosol partitioning of NH₄NO₃ in the seasonality of these components.

Gas-phase NH₃ and aerosol NH₄NO₃ were the dominant species in the total inorganic gas and aerosol species measured in the NEU network. With the current and projected trends in SO₂, NO_x, and NH₃ emissions, concentrations of NH₃ and NH₄NO₃ can be expected to continue to dominate the inorganic pollution load over the next decades, especially NH₃, which is linked to substantial exceedances of ecological thresholds across Europe. The shift from (NH₄)₂SO₄ to an atmosphere more abundant in NH₄NO₃ is expected to maintain a larger fraction of reactive N in the gas phase by partitioning to NH₃ and HNO₃ in warm weather, while NH₄NO₃ continues to contribute to exceedances of air quality limits for PM_{2.5}.

1 Introduction

Air quality policies and research on atmospheric sulfur (S) and nitrogen (N) pollutant impacts on ecosystems and human health have focused on the emissions, concentrations, and depositions of sulfur dioxide (SO₂), nitrogen

oxides (NO_x), ammonia (NH₃), and their secondary inorganic aerosols (SIAs; ammonium sulfate, (NH₄)₂SO₄; ammonium nitrate, NH₄NO₃) (ROTAP, 2012; EMEP, 2019). The aerosols, formed through neutralization reactions between the alkaline NH₃ gas and acids generated in the atmosphere by the oxidation of SO₂ and NO_x (Huntzicker et al., 1980; AQEG, 2012), are a major component of fine particulate matter (PM_{2.5}) (AQEG, 2012; Vieno et al., 2016a) and precipitation (ROTAP, 2012; EMEP, 2019). The negative effects of these pollutants on sensitive ecosystems are mainly through acidification (excess acidity) and eutrophication (excess nutrient N) processes that can lead to a loss of key species and decline in biodiversity (e.g. Hallsworth et al., 2010; Stevens et al., 2010). They are also implicated in radiative forcing and influence climate change through inputs of nitrogen that can alter the carbon cycle (Reis et al., 2012; Sutton et al., 2013; Zaehle and Dalmonech, 2011).

A number of EU policy measures (e.g. 2008/50/EC Ambient Air Quality Directive: EU, 2008; 2016/2284/EU National Emissions Ceilings Directive, NECD: EU, 2016) and wider international agreements (e.g. Gothenburg Protocol; UNECE, 2012) are targeted at abating the emissions and environmental impacts of SO₂, NO_x, and NH₃. The largest emissions reductions have been achieved for SO₂, which decreased by 82 % across the 33 member countries of the European Environment Agency (EEA-33) since 1990, to 4743 kt SO₂ in 2017 (EEA, 2019). Reductions in NO_x emissions have been more modest, at 45 % over the same period, with emissions in 2017 of 8563 kt NO_x exceeding those of SO₂. By contrast, the reductions in NH₃ emissions (of which over 90 % come from agriculture) have been more modest, decreasing by only 18 %. Here, the decrease was largely driven by reductions in fertilizer use and livestock numbers, in particular from eastern European countries, rather than through implementation of any abatement or mitigation measures. More worryingly, the decreasing trend has reversed in recent years, with emissions increasing by 5 % since 2010, to 4788 kt NH₃ in 2017 (EEA, 2019).

In recent assessments, critical loads of acidity were exceeded in about 5 % of the ecosystem area across Europe in 2017 (EMEP, 2018). While the substantial decline in SO₂ emissions has allowed the recovery of ecosystems from acid rain, NH₃ from agriculture and NO_x from transport are increasingly contributing to a larger fraction of the acidity load. Although NH₃ is not an acid gas, nitrification of NH₃ and ammonium (NH₄⁺) releases hydrogen ions (H⁺) that acidify soils and fresh water. The deposition of reactive N (N_r; including oxidized N: NO_x, HNO₃, NO₃⁻, and reduced N: NH₃, NH₄⁺) and its contribution to eutrophication effects have also been identified by the European Environment Agency (EEA) as the most important impact of air pollutants on ecosystems and biodiversity (EEA, 2019). The deposition of N_r throughout Europe remains substantially larger than the level needed to protect ecosystems, with critical-load thresholds for eutrophication from N exceeded in around 62 % of

the EU-28 ecosystem area and in almost all countries in Europe in 2017 (EMEP, 2018).

Following emission, atmospheric transport and fate of the gases are controlled by the following processes: short-range dispersion and deposition, chemical reaction and formation of NH₄⁺ aerosols, and the long-range transport and deposition of the aerosols (Sutton et al., 1998; ROTAP, 2012). Atmospheric S and N_r inputs from the atmosphere to the biosphere occur through (i) dry deposition of gases and aerosols, (ii) wet deposition in rain, and (iii) occult deposition in fog and cloud (Smith et al., 2000; ROTAP, 2012). The deposition processes contribute very different fractions of the total S or N_r input and different chemical forms of the pollutants at different spatial scales. NH₃ is a highly reactive, water-soluble gas and deposits much faster than NO_x (which is not very water-soluble and has low deposition velocity). Dry N deposition by NH₃ therefore contributes a significant fraction of the total N deposition to receptors close to source areas and will often exert the larger ecological impacts compared with other N pollutants (Cape et al., 2004; Sutton et al., 1998, 2007). Numerous studies have shown that N_r deposition in the vicinity of NH₃ sources is dominated by dry NH₃–N deposition (e.g. Pitcairn et al., 1998; Sheppard et al., 2011), with removal of NH₃ close to a source controlled by physical, chemical, and ecophysiological processes (Flechard et al., 2011; Sutton et al., 2007, 2013). Unlike NO_x, HNO₃ (from oxidation of NO_x) is very water-soluble, while NO₃[−] particles can act as cloud condensation nuclei (CCN) so that they are both scavenged quickly and removed efficiently by precipitation. Since NO_x is inefficiently removed by precipitation, wet deposition of NO_x near a source is small and only becomes important after NO_x has been converted to HNO₃ and NO₃[−].

Because of the large numbers of atmospheric N species and their complex atmospheric chemistry, quantifying the deposition of N_r is hugely complex and is a key source of uncertainty for ecosystem effect assessment (Bobbink et al., 2010; Fowler and Reis, 2007; Schrader et al., 2018; Sutton et al., 2007). Input by dry deposition can be estimated using a combination of measured and/or modelled concentration fields with high-resolution inferential models (e.g. Smith et al., 2000; Flechard et al., 2011) or by making direct flux measurements (e.g. Fowler et al., 2001; Nemitz et al., 2008). Although it is possible to measure N_r deposition directly (e.g. Skiba et al., 2009), the flux measurement techniques are complex and resource-intensive, unsuited to routine measurements at a large number of sites. The “inferential” modelling approach provides a direct estimation of deposition from N_r measurements by applying a land-use-dependent deposition velocity (*V_d*) to measured concentrations (Dore et al., 2015; Flechard et al., 2011; Simpson et al., 2006; Smith et al., 2000).

At present, there are limited atmospheric measurements that speciate the gas- and aerosol-phase components at multiple sites over several years. On a European scale, atmo-

spheric measurements of sulfur (SO₂, particulate SO₄^{2−}) and nitrogen (NH₃, HNO₃, particulate NH₄⁺, NO₃[−]) have been made by a daily filter pack method across the European Monitoring and Evaluation Program (EMEP) networks since 1985, providing data for evaluating wet- and dry-deposition models (EMEP, 2016; Tørseth et al., 2012). The method, however, does not distinguish between the gas- and aerosol-phase N species. Consequently, these data are reported as total inorganic ammonium (TIA = sum of NH₃ and NH₄⁺) and total inorganic nitrate (TIN = sum HNO₃ and NO₃[−]), limiting the usefulness of the data. Speciated measurements by an expensive and labour-intensive daily annular denuder method are also made (Tørseth et al., 2012) but are necessarily restricted to a small number of sites due to the high costs associated with this type of measurement. There are also networks with a focus on specific N components, for example, the national NH₃ monitoring networks in the Netherlands (LML; van Zanten et al., 2017) and in the UK (National Ammonia Monitoring Network, NAMN; Tang et al., 2018a) or compliance monitoring across Europe in the case of SO₂ and NO_x. The UK is unique in having an extensive set of speciated gas and aerosol monitoring data from the Acid Gas and Aerosol Network (AGANet), with measurements from 1999 to the present (Tang et al., 2018b).

In this context, there is an ongoing need for cost-effective, easy-to-operate, time-integrated atmospheric measurement for the respective gas and aerosol phases at sufficient spatial scales. Such data would help to (1) improve estimates of N deposition; (2) contribute to development and validation of long-range transport models, e.g. EMEP (Simpson et al., 2006) and EMEP4UK (Vieno et al., 2014, 2016b); (3) interpret interactions between the gas and aerosol phases; and (4) interpret ecological responses to nitrogen (e.g. ecosystem biodiversity or net carbon exchange). To contribute to this goal, a “three-level” measurement strategy in the EU Framework Programme 6 Integrated Project “NitroEurope” (NEU; <http://www.nitroeurope.ceh.ac.uk/>, last access: 29 July 2020) between 2006 and 2010 delivered a comprehensive integrated assessment of the nitrogen cycle, budgets, and fluxes for a range of European terrestrial ecosystems (Sutton et al., 2007; Skiba et al., 2009). At the most intensive level (Level 3), state-of-the-art instrumentation for high-resolution, continuous measurements at just 13 “flux super sites” provided detailed understanding on atmospheric and chemical processes (Skiba et al., 2009). By contrast, manual methods with a low temporal frequency (monthly) at the basic level (Level 1) provided measurements of N_r components at a large number of sites (> 50 sites) in a cost-efficient way in a pan-European network (Tang et al., 2009). Key species of interest included NH₃, HNO₃, and ammonium aerosols ((NH₄)₂SO₄, NH₄NO₃).

In this paper, we present and discuss 4 years of monthly reactive gas (NH₃, HNO₃, HCl) and aerosol (NH₄⁺, NO₃[−], SO₄^{2−}, Cl[−], Na⁺, Ca²⁺, Mg²⁺) measurements from the

Level 1 network set up under the NEU integrated project (Fig. 2). A harmonized measurement approach with a simple, cost-efficient time-integrated method, applied with high spatial coverage, allowed a comprehensive assessment across Europe. The gas and aerosol network was complemented by 2 years of wet-deposition data made at a subset of the sites (Fig. 3). The intention of the smaller bulk wet-deposition network was two-fold: (i) to provide wet-deposition estimates at DELTA[®] (DENuder for Long-Term Atmospheric sampling) sites that do not already have such measurements on site and (ii) to compare the relative importance of reduced and oxidized N versus sulfur in the atmospheric pollution load. Measurements across the network were coordinated between multiple European laboratories. The measurement approach and the operations of the networks, including the implementation of annual inter-comparisons to assess comparability between the laboratories, are described. The data are discussed in terms of spatial and temporal variation in concentrations, relative contribution of the inorganic nitrogen and sulfur components to the inorganic pollution load, and changes in atmospheric concentrations of acid gases and their interactions with NH_3 gas and NH_4^+ aerosol.

2 Methods

2.1 NEU Level 1 DELTA[®] network

The NitroEurope (NEU) Level 1 network was operated between November 2006 and December 2010 to deliver the core measurements of reactive nitrogen gases (NH_3 , HNO_3) and aerosols (NH_4^+ , NO_3^-) for the project (Fig. 1). A low-volume denuder filter pack method, the “DENuder for Long-Term Atmospheric sampling” system (DELTA[®]; Sutton et al., 2001a; Tang et al., 2009, 2018b), with time-integrated monthly sampling was used, which made implementation at a large number of sites possible. Other acid gases (SO_2 , HCl) and aerosols (SO_4^{2-} , Cl^- , Na^+ , Ca^{2+} , Mg^{2+}) were also collected at the same time and measured by the DELTA[®] method. DELTA[®] measurements were co-located with all NEU Level 3 sites with advanced flux measurements (Skiba et al., 2009) and with the network of main CarboEurope-IP CO_2 flux monitoring sites (<http://www.carboeurope.org/ceip/>, last access: 5 January 2020) (Flechard et al., 2011, 2020). Two of the UK sites in the NEU DELTA[®] network are existing UK NAMN (Tang et al., 2018a) and AGANet sites (Tang et al., 2018b). These are Auchencorth Moss (UK-Amo) and Bush (UK-EBu), located in southern Scotland. Monthly gas and aerosol data at the two sites, made as part of the UK national networks, were included in the NEU network. NEU network N_r data were used, together with a range of dry-deposition models, to model dry-deposition fluxes (Flechard et al., 2011) and to assess the influence of N_r on the C cycle, potential C sequestration, and the greenhouse gas balance of ecosystems using CO_2 exchange data from the

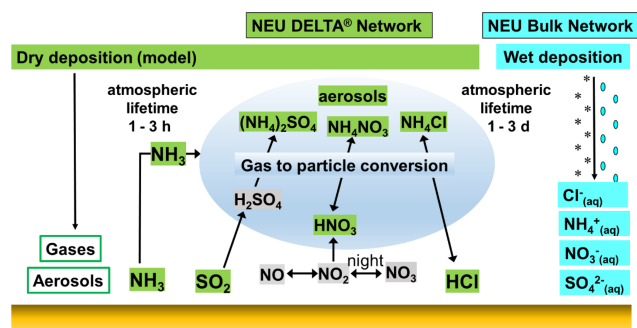


Figure 1. Reaction scheme for the formation of ammonium aerosols from interaction of NH_3 with acid gases HNO_3 , SO_2 , and HCl , showing the components (green) that were measured in the NitroEurope (NEU) DELTA[®] network. Dry deposition of the gas and aerosol components was estimated by inferential modelling (Flechard et al., 2011), while wet deposition (blue) was measured in the NEU bulk wet-deposition network at a subset of the DELTA[®] sites.

co-located CarboEurope sites (Flechard et al., 2020). Other measurements made at the Level 1 sites included estimation of wet-deposition fluxes (Sect. 2.3) and also soil and plant bioassays (Schaufli et al., 2010).

Altogether, the DELTA[®] network covered a wide distribution of sites across 20 countries and four major ecosystem types: crops, grassland, semi-natural, and forests. These sites can be described as “rural” and were chosen to provide a regionally representative estimate of air composition. The network site map is shown in Fig. 2, with site details given in Table S1 in the Supplement. Further information on the sites are also provided in Flechard et al. (2011). Network establishment started in November 2006, with 57 sites operational from March 2007 onwards. Over the course of the network, some sites closed or were relocated due to infrastructure changes, and new sites were also added. A total of 64 sites provided measurements at the end of the project, with 45 of the sites operational the entire time. In addition, replicated DELTA[®] measurements were made at four sites:

1. Auchencorth Moss parallel (P) (UK-AMoP; NH_3 and NH_4^+ measured only);
2. Easter Bush parallel (P) (UK-EBuP; same method as main site);
3. SK04 parallel (P) (SK04P; same method as main site);
4. Fougères parallel (P) (FR-FgsP; different sample train with $2 \times \text{NaCl}$ -coated denuders instead of $2 \times \text{K}_2\text{CO}_3$ -glycerol-coated denuders to capture HNO_3 ; see Sect. 2.2.3) from February to December 2010 only.

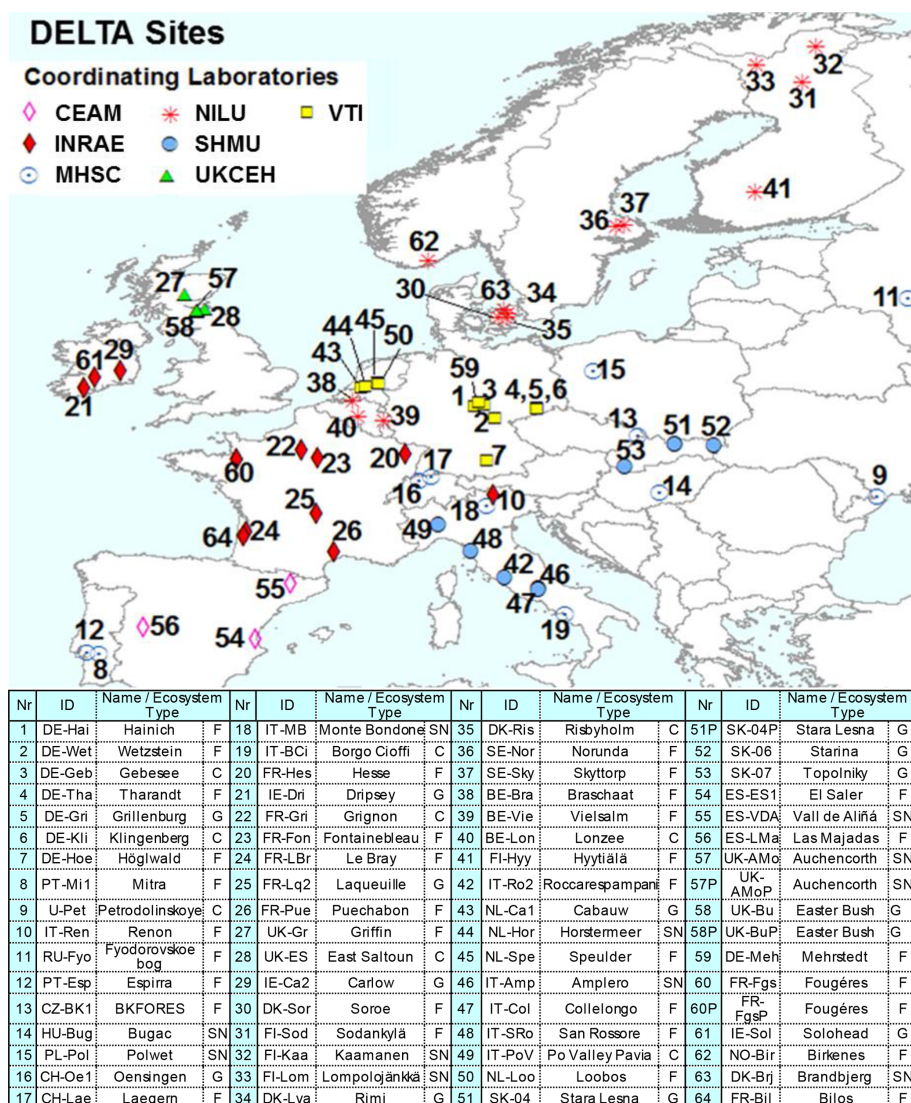


Figure 2. NitroEurope (NEU) DELTA[®] network sites operated between 2006 and 2010. The colour of the symbols indicates the responsible laboratories: CEAM (the Mediterranean Centre for Environmental Studies), vTI (von Thunen Institut), INRAE (French National Research Institute for Agriculture, Food and Environment), MHSC (Meteorological and Hydrological Service of Croatia), UKCEH (UK Centre for Ecology & Hydrology), NILU (Norwegian Institute for Air Research), SHMU (Slovak Hydrometeorological Institute). Ecosystem types are C (crops), G (grassland), F (forests), and SN (short semi-natural; includes moorland, peatland, shrubland, and unimproved and upland grassland). Replicated (P: parallel) DELTA measurements are made at four sites: SK04/SK04P; UK-AMo/UK-AMoP (NH₃ and NH₄⁺ only), UK-Bu/UK-BuP, and FR-Fgs/FR-FgsP (NaCl-coated denuders instead of K₂CO₃–glycerol in sample train).

2.1.1 Coordinating laboratories

A team of seven European laboratories shared responsibility for running the network. Measurement was on a monthly timescale, with each laboratory preparing and analysing the monthly samples with documented analytical methods (see Table S3 for information on analytical methods and limit of detection (LOD)) for between 5 and 16 DELTA sites (Fig. 2). The use of a harmonized DELTA[®] methodology, coupled to defined quality protocols (Tang et al., 2009), ensured comparability of data between the laboratories (see

later in Sect. 3.1 and 3.2). A network of local site operators representing the science teams of each site performed the monthly sample changes and posted the exposed samples back to their designated laboratories for analysis. Air concentration data were submitted by the laboratories for their respective sites in a standard reporting template to UKCEH. Following data checks against defined quality protocols (Tang et al., 2009), the finalized dataset was uploaded to the NEU database (<http://www.nitroeuropa.ceh.ac.uk/>, last access: 29 July 2020). Establishment of the network, including the first year of measurement results on N_r com-

ponents, is reported in Tang et al. (2009). Information on co-located measurements and agricultural activities at each of the sites was also collected and is accessible from the NEU website (<http://www.nitroeuropa.ceh.ac.uk/>, last access: 29 July 2020).

2.2 DELTA[®] methodology

The DELTA[®] method used in the NEU Level 1 network is based on the system developed for the UK Acid Gas and Aerosol Monitoring Network (AGANet, Tang et al., 2018b). Full details of the DELTA[®] method and air concentration calculations in the NEU network are provided by Tang et al. (2009, 2018b). The method uses a small 6 V air pump to deliver low air-sampling rates of between 0.2 and 0.4 L min⁻¹, a high-sensitivity gas meter to record the typical monthly volume of air collected, and a DELTA[®] denuder filter pack sample train to collect separately the gas- and aerosol-phase components. The sample train is made up of two pairs of base- and acid-impregnated denuders (15 and 10 cm long) to collect acid gases and NH₃, respectively, under laminar conditions. A two-stage filter pack with base- and acid-coated cellulose filters collects the aerosol components downstream of the denuders. The base coating used was K₂CO₃–glycerol, which is effective for the simultaneous collection of HNO₃, SO₂, and HCl (Ferm, 1986), while the acid coating was either citric acid for temperate climates or phosphorous acid for Mediterranean climates (Allegrini et al., 1987; Ferm, 1979; Perrino et al., 1990; Fitz, 2002). In this way, artefacts between gas- and aerosol-phase concentrations are minimized (Ferm et al., 1979; Sutton et al., 2001a). The DELTA[®] air inlet has a particle cut-off of ~ 4.5 µm, which means fine-mode aerosols in the PM_{2.5} fraction and some of the coarse-mode aerosols < PM_{4.5} will be collected (Tang et al., 2015).

A low-voltage version of the AGANet DELTA[®] system was built centrally by UKCEH and sent to each of the European sites, where they were installed by local site contacts. These systems operated on either 6 V (off mains power with a transformer) or 12 V from batteries (wind- and solar-powered). Air sampling was direct from the atmosphere without any inlet lines or filters to avoid potential loss of components – in particular HNO₃, which is very “sticky” – to surfaces. Sampling height was 1.5 m above ground or vegetation in open areas. In forested areas, the DELTA[®] equipment was set up either in large clearings or on towers 2–3 m above the canopy (see Flechard et al., 2011).

2.2.1 Calculation of gas and aerosol concentrations

Atmospheric gas and aerosol concentrations in the DELTA[®] method are calculated from the number of inorganic ions (NH₄⁺, NO₃⁻, SO₄²⁻, Cl⁻, and base cations) in the denuder and aerosol aqueous extracts and the volume of air sampled (from gas meter readings), which is typically 15 m³ for a

monthly sample. The volumes of deionized water used to extract acid-coated denuders and aerosols filters are 3 and 4 mL, respectively. For the base-coated denuders and aerosol filters, the extract volume in both cases is 5 mL. An example is shown here for calculating the atmospheric concentrations of NH₃ (gas) (Eq. 1) and NH₄⁺ (aerosol) (Eq. 2) from the aqueous extracts, based on an air volume of 15 m³ collected in a typical month.

$$\text{Gas NH}_3 (\mu\text{g m}^{-3}) = \frac{\text{NH}_4^+ (\text{mg L}^{-1}) [\text{sample} - \text{blank}] \cdot 3 \text{ mL} \cdot \left(\frac{17}{18}\right)}{15 \text{ m}^3} \quad (1)$$

$$\text{Particle NH}_4^+ (\mu\text{g m}^{-3}) = \frac{\text{NH}_4^+ (\text{mg L}^{-1}) [\text{sample} - \text{blank}] \cdot 4 \text{ mL}}{15 \text{ m}^3} \quad (2)$$

Pairs of base- and acid-coated denuders are used to collect the acid gases and alkaline NH₃ gas, respectively. This allows denuder collection efficiency of, for example, NH₃ (Eq. 3) to be assessed as part of the data quality assessment process. An imperfect acid coating on the denuders for example can lead to lower capture efficiencies (Sutton et al., 2001a; Tang and Sutton, 2003).

$$\begin{aligned} \text{Denuder collection efficiency, NH}_3 (\%) \\ = 100 \times \frac{\text{NH}_3 (\text{Denuder 1})}{\text{NH}_3 (\text{Denuder 1} + \text{Denuder 2})} \end{aligned} \quad (3)$$

A correction, based on the collection efficiency, is applied to provide a corrected air concentration (χ_a (corrected); Eq. 4) (Sutton et al., 2001a; Tang et al., 2018a, 2018b). With a collection efficiency of 95 %, the correction amounts to 0.3 % of the corrected air concentration. For an efficiency below 60 %, the correction amounts to more than 50 % and is not applied. The air concentration (χ_a) of NH₃ is then determined as the sum of NH₃ in denuders 1 and 2 (Tang et al., 2018a). By applying the infinite series correction, the assumption is that any NH₃ (and other gases) that is not captured by the denuders will be collected on the downstream aerosol filter. To avoid double-counting, the estimated amount of “NH₃ breakthrough” is subtracted from the NH₄⁺ concentrations on the aerosol filter.

$$\chi_a (\text{corrected}) = \chi_a (\text{Denuder 1}) \cdot \frac{1}{1 - \left[\frac{\chi_a (\text{Denuder 2})}{\chi_a (\text{Denuder 1})} \right]} \quad (4)$$

2.2.2 Estimating sea salt and non-sea-salt SO₄²⁻ (ss-SO₄²⁻ and nss-SO₄²⁻)

Sea salt SO₄²⁻ (ss-SO₄²⁻) in aerosol was estimated according to Eq. (5), based on the ratio of the mass concentrations of

SO₄^{2−} to the reference Na⁺ species in seawater (Keene et al., 1986; O'Dowd and de Leeuw, 2007).

$$[\text{ss} - \text{SO}_4^{2-}] (\mu\text{g ss} - \text{SO}_4^{2-} \text{ m}^{-3}) = 0.25 \cdot [\text{Na}^+] (\mu\text{g Na}^+ \text{ m}^{-3}) \quad (5)$$

Non-sea-salt SO₄^{2−} (nss-SO₄^{2−}) was then derived as the difference between total measured SO₄^{2−} and ss-SO₄^{2−} (Eq. 6).

$$[\text{nss} - \text{SO}_4^{2-}] (\mu\text{g ss} - \text{SO}_4^{2-} \text{ m}^{-3}) = [\text{SO}_4^{2-}] (\mu\text{g SO}_4^{2-} \text{ m}^{-3}) - [\text{ss_SO}_4^{2-}] (\mu\text{g ss} - \text{SO}_4^{2-} \text{ m}^{-3}) \quad (6)$$

2.2.3 Artefact in HNO₃ determination

Results from the first DELTA[®] inter-comparison in the NEU network (Tang et al., 2009) (see also Sect. 2.5) and further work by Tang et al. (2015, 2018b) have shown that HNO₃ concentrations may be overestimated on the carbonate-coated denuders used due to co-collection of other oxidized nitrogen components, most likely from nitrous acid (HONO). In the UK AGANet, HNO₃ data are corrected with an empirical factor of 0.45 derived by Tang et al. (2015). Since the correction factor for HNO₃ is uncertain (estimated to be ±30 %) and derived for UK conditions, no attempt has been made to correct the HNO₃ data from the NEU network. The DELTA[®] method remained unchanged throughout the entire network operation and provided a consistent set of measurements by the same protocol. The caveat is that the HNO₃ data presented in this paper also include an unknown fraction of oxidized N, most probably HONO, and therefore represent an upper limit in the determination of HNO₃. Contribution from NO₂ is likely to be small since this is collected with a low efficiency on carbonate-coated denuders (Bai et al., 2003; Tang et al., 2015), and the network sites are rural, where NO_x concentrations are expected to be in the low parts per billion. At the French Fougères parallel site (FR-FgsP), NaCl-coated denuders were used to measure HNO₃ to compare with results from K₂CO₃–glycerol-coated denuders at the main site (FR-Fgs) (see Sect. 2.1 for methodology and Sect. 3.3.1 for data inter-comparison).

2.3 NEU bulk wet-deposition network

The NEU bulk wet-deposition network (Fig. 3, Table S2) was established to provide wet-deposition data on NH₄⁺ and NO₃[−]. It was set up 2 years after the establishment of the NEU DELTA[®] network, with sites located at a subset of DELTA[®] sites that did not already have on-site wet-deposition measurements. Sampling commenced at some sites in January 2008, with 14 sites operational from March 2008. Site changes also occurred during the operation of this network, again with some site closures and new site additions over time. In total, 12 sites provided 2 years of monthly

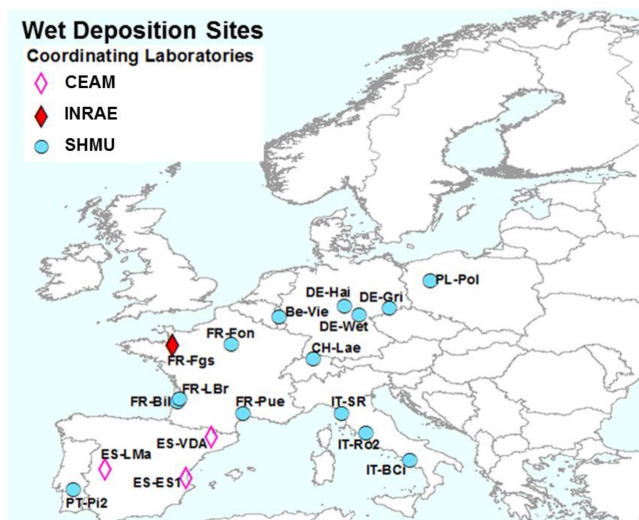


Figure 3. NitroEurope (NEU) bulk wet-deposition network sites operated between 2008 and 2010. The colour of the symbols indicates the responsible laboratories: CEAM (the Mediterranean Centre for Environmental Studies), INRAE (French National Research Institute for Agriculture, Food and Environment), and SHMU (Slovak Hydrometeorological Institute).

data, with a further 6 sites providing 1 year of monthly data between 2008 and October 2010, when measurements ended.

The type of bulk precipitation collector used was a Rotenkamp sampler (Dämmgen et al., 2005), mounted 1.5 m above ground, or in the case of forest sites, either in clearings or above the canopy. Each unit has two collectors providing replicated samples, comprising a pyrex glass funnel (aperture area = 84.9 cm²) with vertical sides, connected directly to a 3 L collection bottle (material: low-density polyethylene), which was changed monthly. Thymol (5-methyl-2-(1-methylethyl)phenol) (150 mg) was added as a biocide (Cape et al., 2012) to a clean, dry pre-weighed bottle at the start of each collection period. This provided a minimum thymol concentration of 50 mg L^{−1} for a full bottle to preserve the sample against biological degradation of labile nitrogen compounds during the month-long sampling.

Three European laboratories shared management and chemical analysis for the network (Fig. 3). The laboratories were CEAM (all three Spanish sites); INRAE (French Renon site); and SHMU, designated the main laboratory responsible for all other sites. A full suite of precipitation chemistry analyses were carried out that included pH, conductivity, NH₄⁺, NO₃[−], SO₄^{2−}, PO₄^{3−}, Cl[−], Na⁺, K⁺, Ca²⁺, and Mg²⁺. Rain volumes and precipitation chemistry data were submitted in a standard template to UKCEH for checking and then uploaded to the NEU database (<http://www.nitroeuropa.ceh.ac.uk/>, last access: 29 July 2020). Samples with high P (> 1 μg L^{−1} PO₄^{3−}), high K⁺, and/or NH₄⁺ values that are indicative of bird contamination were rejected. Annual wet deposition (e.g. kg N ha^{−1} yr^{−1}) was estimated from the product of the

species concentrations and rain volume. Determinations of organic N were also carried out on some of the rain samples in a separate investigation reported by Cape et al. (2012).

2.4 Laboratory inter-comparisons: chemical analysis

All laboratories in the DELTA[®] and bulk wet-deposition networks participated in water chemistry proficiency testing (PT) schemes in their own countries as well as the EMEP (once annually; <http://www.emep.int>, last access: 8 February 2010) and/or WMO-GAW (twice annually; <http://www.qasac-americas.org>, last access: 12 January 2020) laboratory inter-comparison schemes. PT samples for analysis are synthetic precipitation samples for determination of pH, conductivity, and all the major inorganic ions at trace levels. In addition, UKCEH also organized an annual PT scheme for the duration of the project (NEU-PT) to compare laboratory performance in the analysis of inorganic ions at higher concentrations relevant for DELTA[®] measurements. This comprised the distribution of reference solutions containing known concentrations of ions that were analysed by the laboratories as part of their routine analytical procedures.

2.5 Field inter-comparisons: DELTA measurements

Prior to the NEU DELTA[®] network establishment, a workshop was held to provide training to participating laboratories on sample preparation and analysis. This was followed by a 4-month inter-comparison exercise (July to October 2006) between six laboratories at four test sites (Montelibretti, Italy; Braunschweig, Germany; Paterna, Spain, and Auchencorth, UK). Results of the inter-comparison on N_r components were reported by Tang et al. (2009), which demonstrated good agreement under contrasting climatic conditions and atmospheric concentrations of the N_r gases and aerosols. The first DELTA[®] inter-comparison allowed the new laboratories to gain experience in making measurements and was an extremely useful exercise to check how the whole system works, starting with coating of denuders and filters and DELTA[®] train preparation, sample exchange via post, sample handling, and inter-comparing laboratory analytical performance. Further DELTA[®] inter-comparisons between laboratories were conducted each year for the duration of the project, details of which are summarized in Table 1. At each test site, DELTA[®] systems were randomly assigned to each of the participating laboratories. All laboratories provided DELTA[®] sampling trains for each of the inter-comparison sites and carried out chemical analysis on the returned exposed samples. Measurement results were returned in a standard template to UKCEH, the central coordinating laboratory for collation and analysis.

2.6 European emissions data

With the exception of Russia and Ukraine, official reported national emissions data on SO₂, NO_x, and NH₃ are avail-

able for all other 18 countries in the NEU network from the European Environment Agency (EEA) website (EEA, 2020). Emissions data for the period 2007 to 2010 were extracted, and the emission densities of each gas (t km⁻² yr⁻¹) in each country were derived by dividing the 4-year-averaged total emissions by the land area (km²). Gridded emissions data (at 0.1° × 0.1° resolution) for SO₂, NO_x, and NH₃ are available from the EMEP emissions database (EMEP, 2020). The 0.1° × 0.1° gridded data for the period 2007 to 2010 were downloaded and were used to estimate national total emissions (sum of all grid squares in each country) and 4-year-averaged emission densities (t km⁻² yr⁻¹) for Russia and Ukraine. As a check, total emissions for the other 18 countries were also calculated by this method and were the same as the national emission totals reported by the EEA (EEA, 2019).

2.7 National air quality network data from the Netherlands and UK

2.7.1 Dutch LML network data

Atmospheric NH₃ has been monitored at eight sites in the Dutch national air quality monitoring network (LML, Landelijk Meetnet Luchtkwaliteit) since 1993 (van Zanten et al., 2017). The low-density, high-time-resolution LML network is complemented by a high-density monthly diffusion tube network, the Measuring Ammonia in Nature (MAN) network (<http://man.rivm.nl>, last access: 6 November 2018) (Lolkema et al., 2015). The MAN network has 136 monitoring locations sited within nature reserves and includes 60 Natura 2000 sites, with concentrations ranging between 1.0 and 14 µg m⁻³ (Lolkema et al., 2015). The focus of the MAN network is to provide site-based NH₃ concentrations for the nature conservation sites rather than a representative spatial-concentration field for the country. Hourly NH₃ and SO₂ data which were also available from the eight sites in the LML network were downloaded from the Rijksinstituut voor Volksgezondheid en Milieu (RIVM, the Dutch National Institute for Public Health and the Environment) website (<http://www.lml.rivm.nl/gevalideerd/index.php>, last access: 6 November 2018). The 4-year-averaged NH₃ and SO₂ concentrations for the period 2007 to 2010 were calculated and used to complement measurement data from the four Dutch sites in the NEU DELTA[®] network.

2.7.2 UK NAMN and AGANet network data

Atmospheric NH₃, acid gases, and aerosols are measured in the UK NAMN (since 1996) and AGANet (since 1999) (Tang et al., 2018a, b). The UK approach is a high-density network with low-time-resolution (monthly) measurements, combining an implementation of the DELTA[®] method used in the present NEU DELTA[®] network and a passive ALPHA[®] method (Tang et al., 2001) to increase network coverage

Table 1. Details of annual NitroEurope (NEU) DELTA[®] field inter-comparisons conducted between 2006 and 2010.

Inter-comparison period	Test sites	Participating laboratories	Number of monthly measurement periods
2006 (Jul–Oct)	Auchencorth, UK Braunschweig, Germany Montelibretti, Italy Paterna, Spain	6	4
2007 (Jul–Aug)	Auchencorth, UK Montelibretti, Italy	6	2
2008 (Apr–May)	Auchencorth, UK Braunschweig, Germany	7 (INRAE: new laboratory)	2
2009 (Nov–Dec)	Auchencorth, UK Montelibretti, Italy	7 (INRAE: new laboratory)	2

in NH₃ measurements (Sutton et al., 2001b; Tang et al., 2018a). Monthly and annual data for the overlapping period of the project were extracted from the UK-AIR website (<https://uk-air.defra.gov.uk/>, last access: 25 November 2019) and nested with the NEU network data for analysis in this paper.

3 Results and discussion

3.1 Laboratory inter-comparison results: chemical analysis

Figure 4 compares the percentage deviation of results from reference solution concentrations (“true value”) reported by the laboratories for different chemical components in the EMEP, WMO-GAW, and NEU proficiency testing (PT) schemes, combined from 2006 to 2010. Each data point is colour-coded in the graphs according to the laboratory providing the measurements.

Altogether, results from the combined PT schemes produced > 100 observations for each reported chemical component over the 4-year period. The performances of laboratories in Fig. 4 can be summarized in terms of the percentage of reported results agreeing within 10 % of the true values (see summary table below Fig. 4), where the true values represent the nominal concentrations in the aqueous test solutions. The best agreements were for SO₄^{2−} and NO₃[−], with an average of 92 % and 87 % of all reported results agreeing within 10 % of the true value across the concentration range covered in the PT schemes. In the case of NH₄⁺, while an average of 90 % of reported results were within 10 % of the reference at 1 mg L^{−1} NH₄⁺, laboratory performance was poorer (68 % agreeing within 10 %) at lower concentrations (0.1–0.9 mg L^{−1}). Poorer performance at the low concentrations was largely due to two laboratories (CEAM and SHMU), with > 50 % of their results reading high. For Na⁺ and Cl[−], the percentages of results agreeing within 10 % of the refer-

ence were 81 % and 86 %, respectively, across the full range of PT concentrations. At concentrations above 1 mg L^{−1}, the agreement improved and increased to 89 % for Na⁺ and 96 % for Cl[−]. A larger spread around the reference values was provided for the base cations Ca²⁺ and Mg²⁺ at low concentrations (< 1 mg L^{−1}). The percentage of results passing at low concentrations below 1 mg L^{−1} was 36 % (Ca²⁺) and 59 % (Mg²⁺), increasing to 80 % (Ca²⁺) and 90 % (Mg²⁺) above 1 mg L^{−1}. The larger scatter at low concentrations is likely due to uncertainty in the chemical analysis at or close to the method limit of detection and reflects challenges of measuring base cations, in particular Ca²⁺ as this is very “sticky” and adsorbs–desorbs from surfaces, leading to analytical artefacts.

To show what the PT reference solution concentrations would correspond to if they were a denuder and/or aerosol extract, equivalent gas (Eq. 1) and/or aerosol concentrations (Eq. 2) (Sect. 2.2.1) are calculated for each of the ions and provided in the summary table in Fig. 4. A 0.5 mg L^{−1} NH₄⁺ solution, for example, is equivalent to an atmospheric concentration of 0.09 µg NH₃ m^{−3} (gas) or 0.13 µg NH₄⁺ m^{−3} (aerosol) for a monthly sample. In Fig. 5, scatterplots are shown comparing all NEU laboratory-reported results with PT reference, where all ion concentrations (mg L^{−1}) from Fig. 4 have been converted to equivalent gas and aerosol concentrations (µg m^{−3}), based on a typical volume of 15 m³ over a month. With the exception of a small number of outliers, most data points are close to the 1 : 1 line, with laboratory results agreeing within ±0.05 µg m^{−3} in equivalent gas and/or aerosol concentrations. These are low ambient concentrations and show that the measurement uncertainty in the analysis of very low concentrations in the PT schemes will be small for the majority of sites in the network, where concentrations were found to be much higher (see Fig. 6).

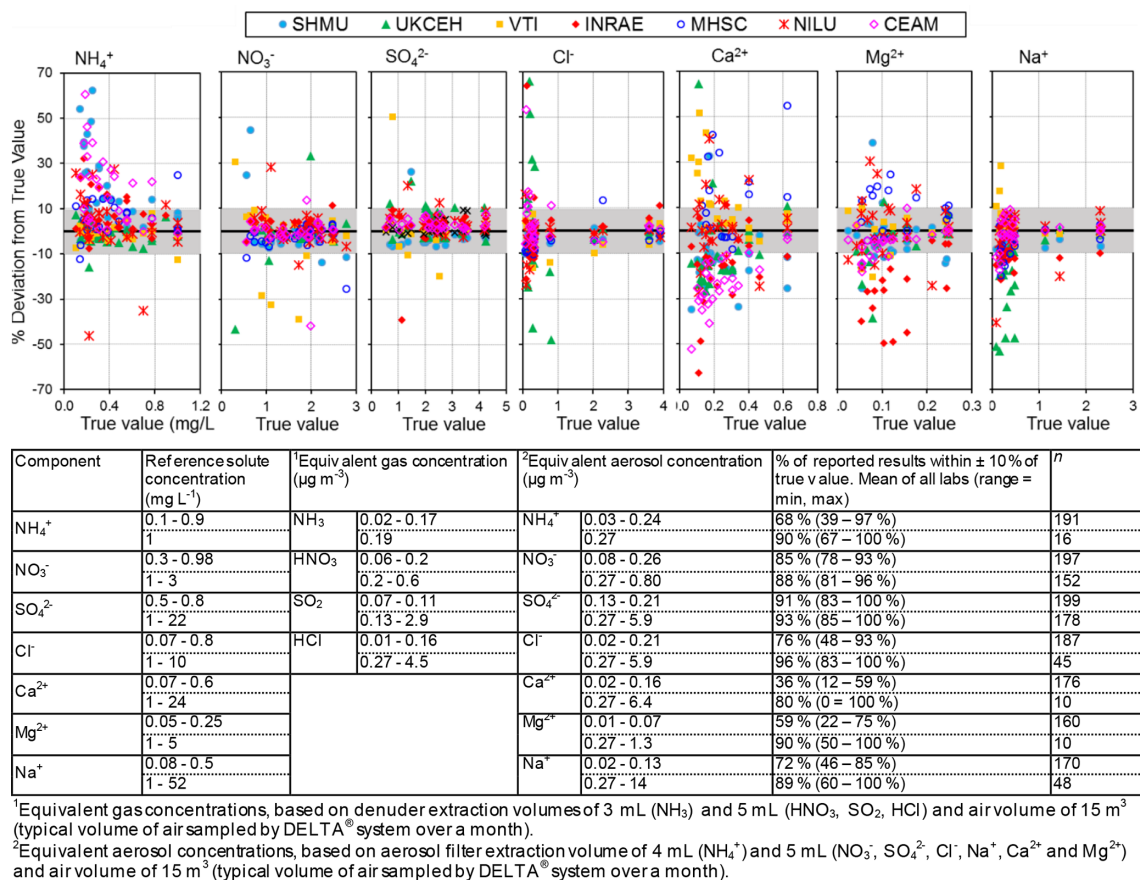


Figure 4. Summary of reported results from all laboratories in chemistry proficiency testing (PT) schemes for chemical analysis of aqueous inorganic ions (2006–2010: EMEP, WMO-GAW, and NitroEurope), expressed as a percentage deviation from the true value (PT reference solutions). The grey shaded areas in the graphs show values that are within $\pm 10\%$ of true value.

3.2 Field inter-comparison results: DELTA[®] measurements

Results from 4 years of annual DELTA[®] field inter-comparisons (2006–2009), for all field sites, are combined and summarized in Fig. 6. The gas and aerosol concentrations measured and reported by each of the laboratories are compared with the median estimate of all laboratories in each of the scatterplots, with the colour of the symbols identifying the laboratory providing the measurements. Regression results (slope and R^2) in the table below the plots provide the main features of the inter-comparison. The slope is equivalent to the mean ratio of each laboratory against the median value, where values close to unity indicate closer agreement to the median value. Overall, the scatterplots show good agreement between the laboratories, with some laboratories showing very close agreement to the median estimates and more scatter observed from the others.

The occurrence of outliers in some of the individual monthly values indicates that caution needs to be exercised in the interpretation of these data points in the inter-comparison. To average out the influence of a few individ-

ual outliers, the mean concentrations from each of the seven laboratories for each of the four field sites were calculated and compared with averaged median estimates of all laboratories for each site. A summary of the mean concentrations and the percentage difference from the median is presented in Table 2. Since the INRAE laboratory did not join the NEU network until 2008, averaged median values from the 2008 and 2009 inter-comparisons are used to compare with the INRAE results, included in the table for clarity. The mean concentrations between laboratories are broadly comparable. Each of the laboratories were also able to resolve the main differences in mean concentrations at the four field sites, ranging from the lowest concentrations at Auchen-corth (e.g. median = $1.4 \mu\text{g NH}_3 \text{ m}^{-3}$) to higher concentrations, representing a more polluted site at Paterna (e.g. median = $5.2 \mu\text{g NH}_3 \text{ m}^{-3}$) for the test periods (Table 2). Larger differences for HCl, Ca²⁺, and Mg²⁺ are due to clear outliers from one or two laboratories at the very low concentrations of these species encountered and may be related to measurement uncertainties at the low air concentrations. The compa-

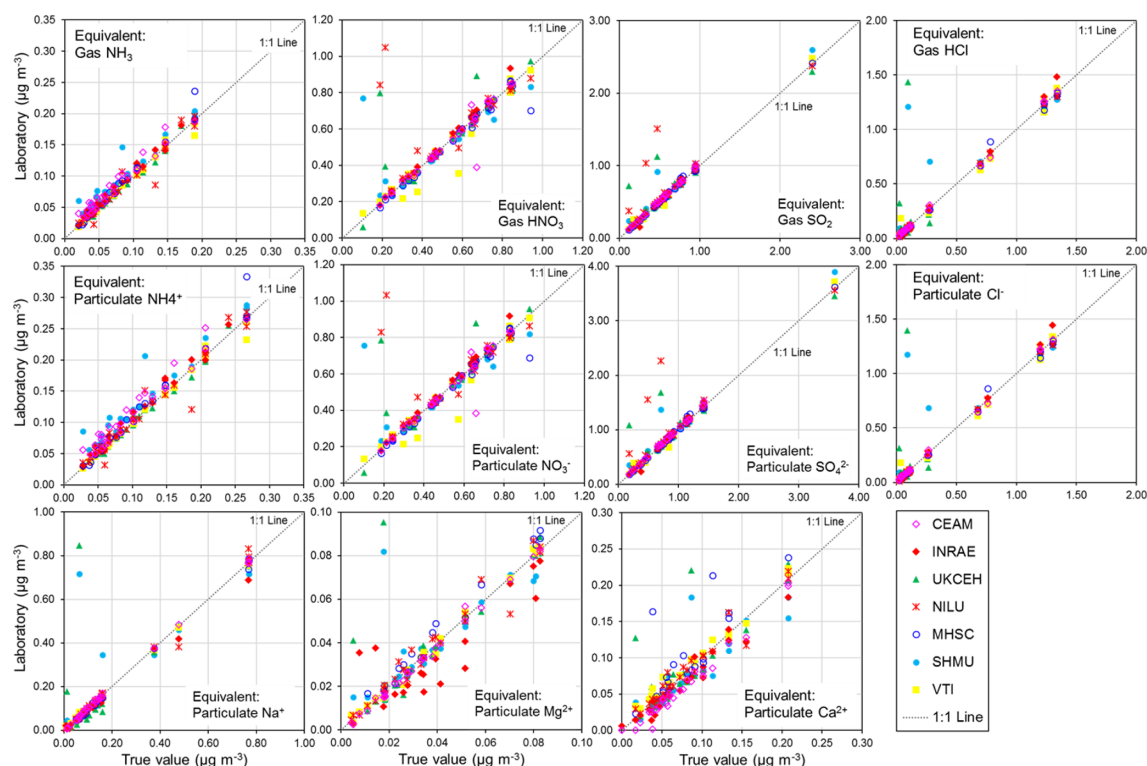


Figure 5. Scatterplots comparing all NEU laboratory-reported results from wet chemistry proficiency testing (PT) schemes (2006–2010: EMEP, WMO-GAW, and NitroEurope) vs. true values (PT reference solutions). All aqueous ion concentrations (mg L^{-1}) from Fig. 4 are converted to equivalent gas and aerosol concentrations ($\mu\text{g m}^{-3}$) for the comparisons.

rability between laboratories for each of the components is next considered in turn.

3.2.1 Inter-comparisons: NH_3 , NH_4^+ , HNO_3 , NO_3^-

The best agreement between laboratories was for the N_r gases (NH_3 , HNO_3) and aerosol species (NH_4^+ , NO_3^-), with slopes within $\pm 10\%$ of the median values and $R^2 > 0.9$ in the regression analysis from five of the laboratories (Fig. 6, Table 2). This is important since N_r species were the primary focus for the NEU DELTA[®] network. Slightly poorer agreement for NH_3 and NH_4^+ was provided by CEAM and MHSC laboratories, with data points both above and below the 1 : 1 line (Fig. 6). The outliers above the 1 : 1 line from MHSC were from the 2006 inter-comparison exercise. Removal of these 2006 outliers improved the MHSC regression slope for NH_3 from 1.21 ($R^2 = 0.87$, $n = 41$) to 0.99 ($R^2 = 0.99$, $n = 10$) (Fig. S1). While this seems to suggest that the performance of MHSC for NH_3 improved following the first inter-comparison exercise, the regression slope for aerosol NH_4^+ increased instead from a slope of 1.26 ($R^2 = 0.83$, $n = 41$) to 1.48 ($R^2 = 0.93$, $n = 10$), suggesting an overestimation of NH_4^+ concentrations (Fig. S1). A possible cause may be the quality and/or variability in the aerosol filter blank values for NH_4^+ as laboratory blanks are subtracted from exposed samples to estimate aerosol NH_4^+ concentrations. While the

laboratory blanks reported by MHSC for aerosol NH_4^+ were low (mean = $0.48 \mu\text{g NH}_4^+$) and smaller than other laboratories (mean = 0.64 – $1.20 \mu\text{g NH}_4^+$) (Fig. S2), their field blanks in the 2006 DELTA inter-comparison exercise were on average 5.5 times larger than the laboratory blanks. This is likely due to extensive delays in getting samples released from customs in Slovakia at the start of the network. Another possibility is a breakthrough of NH_3 from the acid-coated denuders onto the aerosol filters. The denuder collection efficiency of NH_3 gas (Eq. 3, Sect. 2.2.1) reported by MHSC was on average 88 % for all years and 91 % where 2006 data have been excluded (Table S3). This is comparable with the mean collection efficiencies of all laboratories (91 % and 90 %) (Table S4), which makes NH_3 breakthrough an unlikely explanation for the higher readings. The assessment of NH_4^+ is however more uncertain from the reduced number of data points ($n = 10$).

For the CEAM laboratory, reported NH_3 concentrations were on average 16 % lower ($n = 41$) than the median, with a slope of 0.89 ($R^2 = 0.87$), and particulate NH_4^+ was on average 13 % lower ($n = 41$) than the median, with a slope of 0.42 ($R^2 = 0.22$) (Fig. 6). A need to improve the NH_4^+ analysis (indophenol colorimetric assay) in the acid-coated denuders and aerosol filters by the CEAM laboratory was identified from the 2006 inter-comparison (Tang et al., 2009). The

Table 2. Inter-comparison of results from seven European laboratories at four different field test sites for all years (2006–2010). The results shown are the mean concentrations from each laboratory for each site and the averaged median estimates derived from all laboratories for each site.

Site	Median (all years)	CEAM	% diff	UKCEH	% diff	MHSC	% diff	NILU	% diff	SHMU	% diff	vTI (2008/09)	% diff	* Median	*INRAE	*% diff
NH₃																
Auchencorth	1.42	1.23	−13	1.39	−2	1.51	6	1.60	13	1.48	4	1.38	−2	1.06	1.17	10
Braunschweig	4.32	3.61	−16	4.34	0	4.62	7	4.87	13	4.27	−1	4.41	2	6.40	6.64	4
Montelibretti	2.46	1.66	−33	2.44	−1	2.89	18	2.77	12	2.63	7	2.34	−5	1.91	1.91	0
Paterna	5.21	4.39	−16	5.27	1	7.00	34	6.22	19	5.55	7	4.57	−12			
NH₄⁺																
Auchencorth	0.73	0.69	−6	0.64	−13	0.92	26	0.73	0	0.96	31	0.74	2	0.58	0.60	2
Braunschweig	1.55	1.54	−1	1.61	4	2.15	39	1.18	−24	1.64	6	1.45	−6	1.38	1.31	−5
Montelibretti	0.95	0.87	−9	0.86	−9	1.21	27	0.72	−24	1.13	19	0.93	−3	0.96	0.96	0
Paterna	1.80	0.50	−72	1.56	−13	2.12	18	1.64	−9	2.04	13	2.26	25			
HNO₃																
Auchencorth	0.57	0.57	−1	0.53	−7	0.69	21	0.62	9	0.59	3	0.49	−15	0.55	0.59	7
Braunschweig	2.36	1.79	−24	2.82	19	2.67	13	2.43	3	2.48	5	2.09	−11	2.85	2.85	0
Montelibretti	2.64	2.53	−4	2.74	4	3.08	17	2.60	−2	2.77	5	2.31	−13	1.70	1.70	0
Paterna	2.67	2.82	6	2.73	2	3.18	19	2.61	−2	2.40	−10	2.05	−23			
NO₃[−]																
Auchencorth	1.21	1.24	3	1.18	−2	1.16	−4	1.27	4	1.20	−1	1.18	−3	1.26	1.14	−9
Braunschweig	3.26	3.70	14	3.43	5	3.33	2	2.28	−30	3.09	−5	2.36	−28	2.92	2.94	1
Montelibretti	1.81	2.00	10	1.84	1	1.57	−13	1.28	−29	1.91	5	1.56	−14	2.11	2.11	0
Paterna	4.52	4.73	5	4.34	−4	4.60	2	4.34	−4	4.57	1	4.32	−4			
SO₂																
Auchencorth	0.95	0.91	−4	0.88	−7	0.99	4	1.10	15	0.91	−4	1.05	10	0.93	1.21	30
Braunschweig	1.49	1.33	−11	1.49	0	1.65	10	1.32	−12	1.41	−5	1.45	−3	1.05	1.17	11
Montelibretti	1.12	1.29	15	1.15	2	1.48	31	0.94	−16	1.45	29	0.99	−12	0.54	0.54	0
Paterna	1.96	2.07	6	1.96	0	2.04	4	1.93	−2	1.99	2	1.78	−9			
SO₄^{2−}																
Auchencorth	1.04	1.21	17	0.80	−23	1.14	10	1.66	60	1.23	19	0.97	−7	0.82	0.58	−29
Braunschweig	2.04	2.67	31	2.12	4	2.35	15	1.58	−22	1.72	−16	1.51	−26	1.61	1.37	−15
Montelibretti	1.55	1.89	22	1.35	−13	1.61	4	1.49	−4	1.79	16	1.43	−8	0.83	0.83	0
Paterna	3.28	4.19	28	3.06	−7	3.06	−7	3.68	12	3.01	−8	3.21	−2			
HCl																
Auchencorth	0.20	1.01	396	0.19	−9	0.15	−28	0.21	4	0.33	62	0.19	−6	0.22	0.74	244
Braunschweig	0.39	1.35	247	0.22	−43	0.16	−59	0.08	−78	0.63	62	0.35	−9	0.16	0.10	−37
Montelibretti	0.40	1.01	151	0.33	−18	0.40	−1	−	−	0.58	45	0.36	−11	0.54	0.54	0
Paterna	0.73	1.77	141	0.42	−42	0.47	−36	−	−	1.32	80	0.81	10			
Cl[−]																
Auchencorth	0.84	0.93	10	0.73	−13	0.86	3	0.26	−69	1.17	39	0.85	1	0.95	0.81	−15
Braunschweig	0.52	0.78	51	0.35	−32	0.57	10	−	−	0.81	56	0.36	−30	0.33	0.21	−39
Montelibretti	0.85	0.94	11	0.76	−11	0.84	−1	−	−	1.19	41	0.86	1	0.66	0.66	0
Paterna	1.37	1.74	27	1.11	−19	1.31	−5	−	−	2.10	54	1.06	−23			
Na⁺																
Auchencorth	0.53	0.79	47	0.55	2	0.60	13	1.25	134	0.68	28	0.56	5	0.65	0.57	−11
Braunschweig	0.37	0.38	4	0.21	−43	0.37	1	0.24	−34	0.85	131	0.37	1	0.27	0.19	−29
Montelibretti	0.59	0.99	67	0.62	4	0.70	18	−	−	0.84	42	0.59	−1	0.51	0.51	0
Paterna	0.94	−	−	1.01	7	0.71	−25	−	−	0.94	−1	0.95	1			
Ca²⁺																
Auchencorth	0.06	0.06	−5	0.06	−11	0.32	415	0.15	137	0.05	−27	0.06	−12	0.03	0.04	38
Braunschweig	0.16	0.07	−57	0.14	−15	0.61	272	0.36	122	0.09	−47	0.11	−34	0.07	0.08	15
Montelibretti	0.16	0.54	241	0.16	−1	0.45	183	−	−	0.15	−4	0.16	2	0.08	0.08	0
Paterna	0.64	−	−	0.53	−17	1.69	163	−	−	0.49	−24	0.57	−12			
Mg²⁺																
Auchencorth	0.05	0.07	27	0.05	−3	0.14	172	0.18	251	0.05	−6	0.05	−8	0.05	0.09	65
Braunschweig	0.05	0.03	−33	0.04	−26	0.10	114	0.08	61	0.03	−35	0.02	−56	0.02	0.04	77
Montelibretti	0.06	0.13	113	0.06	−2	0.18	185	−	−	0.05	−13	0.06	2	0.04	0.04	0
Paterna	0.13	−	−	0.13	−4	0.33	147	−	−	0.10	−24	0.13	−2			

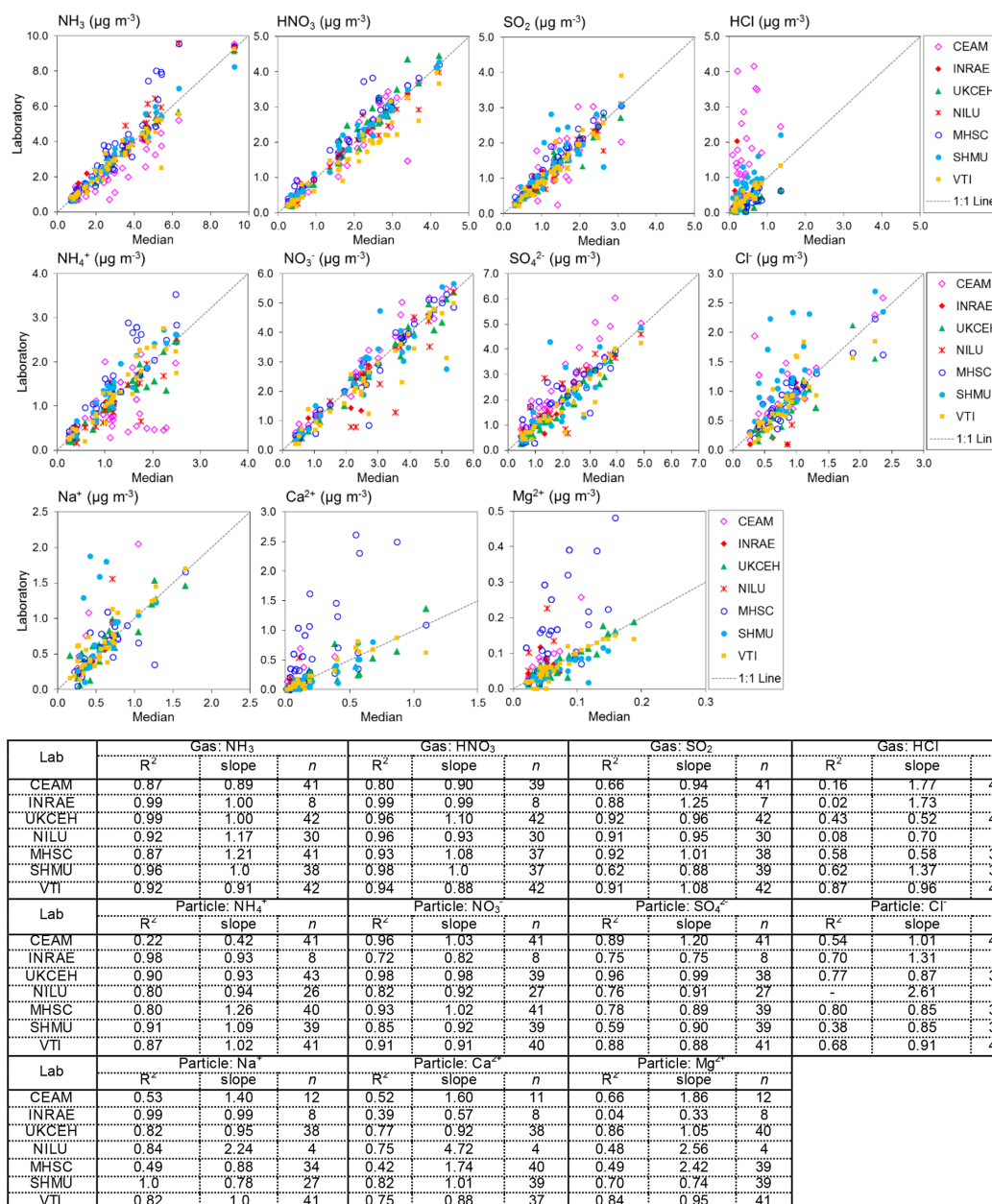


Figure 6. Scatterplots comparing atmospheric gas (NH₃, HNO₃, SO₂, and HCl) and aerosol (NH₄⁺, NO₃⁻, SO₄²⁻, Cl⁻, Na⁺, Ca²⁺, Mg²⁺) concentrations measured by each of the NEU laboratories with the median estimate of all laboratories. Data from all field inter-comparisons (2006–2009) for all test sites (Auchencorth, UK; Braunschweig, Germany; Montelibretti, Italy; and Paterna, Spain) are combined in the analysis. A summary of the regression results is shown in the table below the graphs. Note that (i) there are fewer data points for INRAE because they joined the NEU network later in 2007 and participated in the 2008 and 2009 inter-comparisons only, and (ii) the low number of observations in some cases was due to some laboratories not reporting all parameters. NILU: HCl, Cl⁻, Na⁺, Ca²⁺, and Mg²⁺ reported for 2008 inter-comparisons only; CEAM: Na⁺, Ca²⁺, Mg²⁺ reported for 2007–2009 inter-comparisons only.

indophenol method for aqueous NH₄⁺ determination is pH-sensitive. Calibration solutions and quality control checks for the colorimetric assays are made up in deionized water (pH 7), whereas the aqueous extracts from the DELTA[®] acid-coated denuders and cellulose filters are acidic (pH ~ 3). Determination of NH₄⁺ in the denuder extracts may therefore

be underestimated if the pH of the indophenol reaction has not been adjusted for the increased acidity in the sample extracts. When the 2006 data are excluded from the regression analysis, the slopes for NH₃ and NH₄⁺ increased to 1.02 ($R^2 = 0.94$, $n = 12$) and 0.98 ($R^2 = 0.51$, $n = 12$), respectively (Fig. S1). The improved agreement with other lab-

oratories after the 2006 inter-comparison suggests that the method under-read was largely resolved, reflected in an improvement in the slope. Despite some uncertainties in the NH₃ and NH₄⁺ measurements, the laboratories were able to clearly resolve the main differences in mean concentrations at the four different field sites in all years (Table 2). The results presented here for CEAM and MHSC highlight the importance of the initial inter-comparison exercise in identifying and resolving sampling and analytical issues at the start of the project.

3.2.2 Inter-comparisons: SO₂, SO₄²⁻

Six laboratories provided slopes within 12 % of the median values in the regression analysis for SO₂ (Fig. 6). The smaller R^2 values were from two laboratories (CEAM and SHMU; $R^2 < 0.7$), with data points both above and below the 1 : 1 line. For INRAE, the larger slope of 1.6 ($R^2 = 9$) was due to a single high SO₂ reading reported for Auchencorth of 2.0 µg SO₂ m⁻³, compared with the median of 1.4 µg SO₂ m⁻³. When the mean SO₂ concentrations measured by INRAE are compared with the median, the difference was on average 13 %, providing acceptable agreement, which suggests that the high reading may just be an outlier. There was more scatter in the inter-comparison for SO₄²⁻, although the majority of points are still close to the 1 : 1 line (Fig. 6). Six laboratories provided slopes within 12 % of the median values in the regression analysis also for SO₄²⁻. The regression slope from CEAM for SO₄²⁻ was 1.2 ($R^2 = 0.9$), which is still within 20 % of the median. The SO₂ and SO₄²⁻ measurements were broadly comparable between the laboratories, with mean concentrations agreeing on average within 6 % of the median (Table 2).

3.2.3 Inter-comparisons: HCl, Cl⁻

The HCl inter-comparison shows clear outliers from the CEAM laboratory, with concentrations that were on average up to 2 times higher than other laboratories (slope = 1.8). For example, a mean concentration of 1.8 µg HCl m⁻³ was reported by CEAM for Paterna, compared with a median of 0.7 µg HCl m⁻³. Apart from CEAM, the mean concentrations of HCl reported by the other laboratories were generally comparable (Table 2). The larger percentage differences between the measured mean and median at each site reflect the challenges of measuring the very low concentrations of HCl at these sites of < 0.5 µg HCl m⁻³ (slightly higher at Paterna). HCl results were reported by NILU for the 2008 inter-comparison exercise only, limiting the number of measurements ($n = 4$) available for comparison.

The comparison for Cl⁻ showed better agreement of the CEAM laboratory results with other laboratories in both the inter-comparison of individual monthly values (Fig. 6) and the mean concentrations (Table 2). Like HCl, larger percentage differences between the measured concentrations and

median at each site may be attributed to higher measurement uncertainties at the low concentrations of Cl⁻. For NILU, there were only two data points for Cl⁻ from the Auchencorth site in the 2008 inter-comparison. Overall, the inter-comparison for HCl and Cl⁻ showed that the laboratories were able to resolve the main differences in mean concentrations at the different sites even at the low concentrations encountered.

3.2.4 Inter-comparisons: base cations (Na⁺, Ca²⁺, Mg²⁺)

Measurements of Ca²⁺ and Mg²⁺ were the most uncertain, with the largest scatter in the inter-comparisons (Fig. 6). Despite the trace levels of these base cations at all field sites, four laboratories (INRAE, UKCEH, SHMU, vTI) provided data close to the 1 : 1 line, demonstrating close agreement between these laboratories. The clear outliers above the 1 : 1 line are from CEAM, MHSC, and NILU, with slopes > 2. While MHSC over-read Ca²⁺ and Mg²⁺, its results for Na⁺ were in better agreement with other laboratories, with a slope of 0.9 ($R^2 = 0.5$) (Fig. 6). There was a lot of scatter in the data however, with outlier points both above and below the 1 : 1 line, suggesting measurement uncertainties in their base cation measurements. For NILU, the only base cation results reported by the laboratory were for the 2008 DELTA[®] inter-comparisons at Auchencorth and Braunschweig. This accounts for the low number of data points ($n = 4$) from the NILU laboratory. The median concentrations of Ca²⁺ and Mg²⁺ at both field sites were very low (< 0.1 µg m⁻³), which makes comparison with the few data reported from NILU highly uncertain. Like NILU, CEAM also did not report base cation results for all of the DELTA[®] inter-comparison. Base cation results provided by CEAM were for 2007–2009 only.

3.3 Variation in annual mean gas and aerosol concentrations and composition

3.3.1 Comparisons according to ecosystem types

Annual averaged concentrations of gases and aerosols measured in the NEU DELTA[®] network are presented in Fig. 7, with sites grouped according to each of the four major ecosystem types: crops, grassland, forests, and semi-natural. These are the classifications used in dry-deposition models, where ecosystem-specific deposition velocities (V_d) are combined with measurement data to produce estimates of N_r dry deposition (Flechard et al., 2011).

A total of 64 sites from 20 different countries, including replicated measurements at four of the sites, are compared in Fig. 7. Not all of the sites however were operational all of the time or at the same time. Changes in the numbers and locations of sites occurred over the duration of the network, for example, due to site closures, relocations, and/or new site additions. The annual averaged concentrations plotted for each

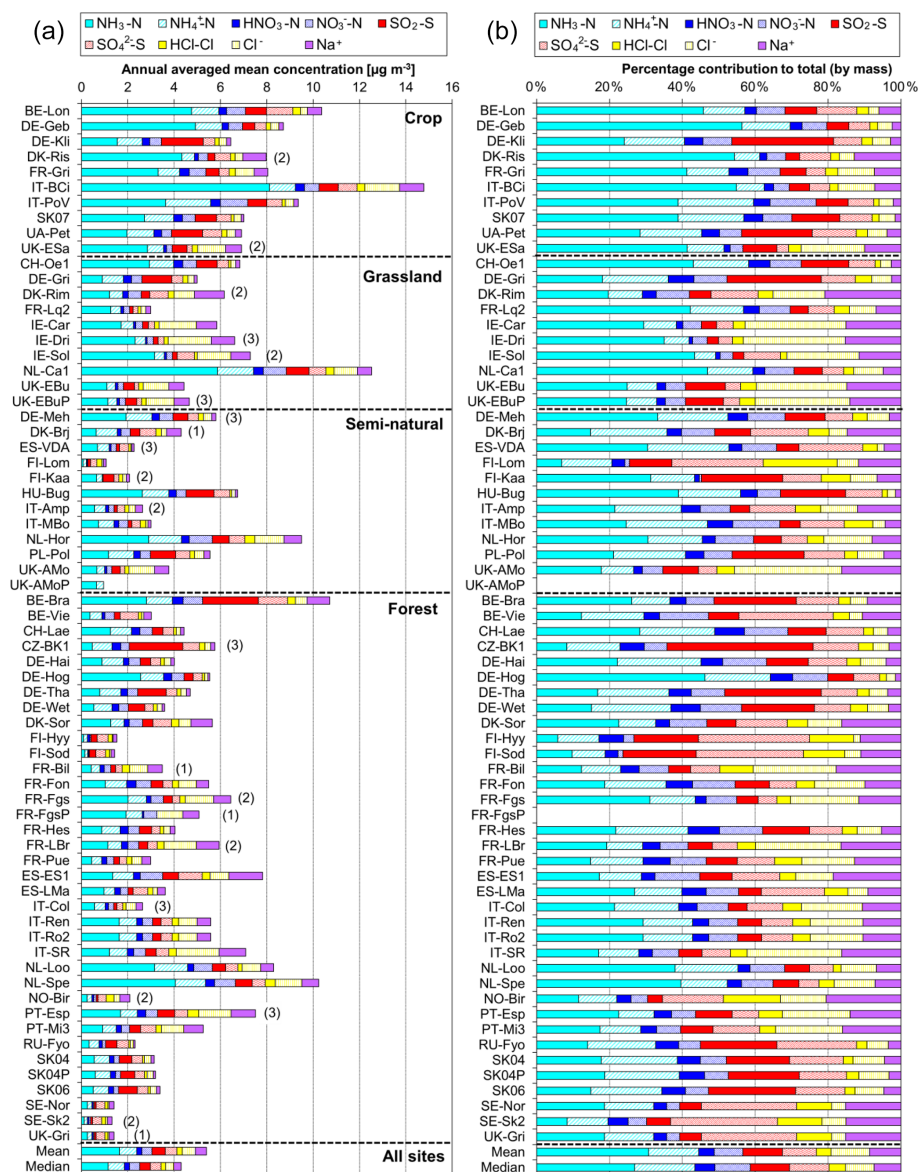


Figure 7. (a) Annual averaged gas and aerosol concentrations (2007–2010) of sites in the NEU DELTA[®] network, grouped according to ecosystem type: crops ($n = 10$), grassland ($n = 9 + 1$ parallel), semi-natural ($n = 11 + 1$ parallel), and forests ($n = 34 + 2$ parallel). (b) Percentage composition of gas and aerosol components measured at NEU DELTA[®] network sites ($n = 64 + 4$ parallel sites) (mean of all annual mean concentrations from 2007 to 2010). Years with < 7 months of data, including 2006, are excluded. Where the number of years contributing to the annual average is < 4 , the number is shown in brackets beside the site data. Ca^{2+} and Mg^{2+} data are not included as these were mostly at or below the limit of detection. Replicated DELTA measurements are made at four sites: FR-Fgs/FR-FgsP (NaCl instead of K_2CO_3 -glycerol-coated denuders; HCl not measured), SK04/SK04P, UK-Ebu/UK-EBuP, and UK-AMo/UK-AMoP (NH_3 and NH_4^+ only).

site are the mean of all available annual means. Where the annual averaged concentration is derived from less than 4 full years of data, the number of years providing the mean is shown, in brackets, next to the site data in the graph. To avoid bias in the calculation of annual means, due to seasonality in the data (see later in Sect. 3.5), years with incomplete data coverage (< 7 months of data in any year) were excluded. Applying these data exclusions, the number of sites that provided annual data was 55 for 2007, 57 for 2008, 54 for 2009,

and 55 for 2010. The number of sites that provided annual data for each year over the entire period was 45 sites.

Sites with parallel (P) DELTA[®] measurements were Auchencorth Moss (UK-AMoP), Easter Bush (UK-EBuP), Fougères (FR-FgsP), and SK04P (EMEP site in Slovakia) (Fig. 7). Overall, good reproducibility in DELTA[®] measurements was demonstrated by the parallel measurements (Figs. S3–S6). At the Auchencorth Moss parallel site (UK-AMoP), NH_3 and NH_4^+ only were measured, and agreement

for these two components was on average within 4 % at the low concentrations measured at this site (annual mean: 0.5–0.9 µg NH₃ m⁻³ and 0.3–0.5 µg NH₄⁺ m⁻³) (Table S5). Parallel measurements at Easter Bush (UK-EBuP) stopped in March 2010. With the exception of Ca²⁺ and Mg²⁺, the comparison of annual mean data from the replicated measurements for 2006 to 2009 provided excellent agreement of 4 % (NO₃⁻) to 12 % (NH₃⁻) at Easter Bush (Table S6). At Fougères (Table S7), HNO₃ concentration measured on K₂CO₃–glycerol-coated denuders (FR-Fgs) was about 2-fold higher than on NaCl-coated denuders in the parallel DELTA[®] system (FR-FgsP), consistent with overestimation of HNO₃ (on average 45 %) on carbonate-coated denuders (see Sect. 2.2.3). The disadvantage of a NaCl coating, however, is that it can only collect HNO₃ and not the other acid gases. A third carbonate denuder is necessary in the sample train to collect and measure SO₂ since SO₂ is only partially captured, and HCl cannot be measured on NaCl denuders (Tang et al., 2015, 2018b). This explains the smaller SO₂ concentrations reported by the FR-FgsP site, with breakthrough of SO₂ (inefficiently captured by NaCl denuders) onto the aerosol filters resulting in larger particulate SO₄²⁻ concentrations than the FR-Fgs site. For the SK04 site, measurement reproducibility for the 4 years of parallel data for the N and S components was good, with agreement ranging from 1.2 % (NH₄⁺) to 9 % (SO₄²⁻) (Table S8). HCl and Na⁺ determinations were however more uncertain, with differences of 67 % and 43 %, respectively (Table S8). It has to be noted, however, that the concentrations of the two components were very low, at < 0.2 µg HCl m⁻³ and < 0.4 µg Na⁺ m⁻³. The differences in concentrations are therefore actually within ± 0.1 µg m⁻³ for HCl and within ± 0.2 µg m⁻³ for Na⁺.

A key feature in Fig. 7 is the dominance of N over S species at most sites, when expressed as micrograms per cubic metre of the element. The mean percentage contribution of sum N_r (NH₃–N, HNO₃–N, NH₄⁺–N, NO₃⁻–N) concentrations to the total mass of gas and aerosol species measured is 52 % (range = 24 %–80 %), twice as much as from sum S (SO₂–S and SO₄²⁻–S; mean = 23 %, range = 7 %–53 %) (Fig. 8). This is consistent with more substantial reductions in SO₂ emissions (–72 %) than achieved with NO_x (–43 %) or NH₃ (–18 %) in Europe between 1991–2010 (EEA, 2019). The differences in atmospheric composition of S and N species in the present assessment therefore reflected changes in emissions of the precursor gases and are also in agreement with a recent assessment of air quality trends showing important changes in S and N composition in air and rain across the EMEP networks (EMEP, 2016).

Most of the N_r concentrations at each site in turn are dominated by reduced N (NH₃–N, NH₄⁺–N) rather than by oxidized N species (HNO₃–N (includes other oxidized N compounds; see Sect. 2.2.3) and NO₃⁻–N). Of the sum N_r concentrations measured, 60 %–97 % (mean = 76 %, *n* = 66) were reduced N (N_{red}) (Fig. 8). Even more strikingly, NH₃ (NH₃–

N) was by far the single most dominant component at the majority of sites, contributing on average 42 % (range = 24 %–56 %, *n* = 10) at cropland sites and 20 % (6 %–46 %, *n* = 35) of the total gas–aerosol concentrations at forest sites (Fig. 8). This illustrates very clearly the importance of NH₃ and by association agricultural emissions in contributing to NH₃–N concentrations and deposition in Europe, with 92 % of total NH₃ emissions in Europe estimated to come from agriculture (EEA, 2019). The reaction of NH₃ with the acid gases HNO₃ and SO₂ forms NH₄⁺-containing particulate matter (PM) that is primarily NH₄NO₃ and (NH₄)₂SO₄ (Fig. 1) (see Sect. 3.4). Together, particulate NH₄⁺–N, NO₃⁻–N and SO₄²⁻–S made up on average 28 % (17 %–40 %, *n* = 10) of the total gas–aerosol concentrations measured at cropland sites (Fig. 8). At semi-natural and forest sites however, that number was even bigger, at 33 % (20 %–40 %, *n* = 11) and 37 % (24 %–57 %, *n* = 35), respectively (Fig. 8).

Secondary NH₄⁺ particles are mainly in the “fine” mode with diameters of less than 2.5 µm (PM_{2.5}) and estimated to contribute between 10 % and 50 % of ambient PM_{2.5} mass concentration in some parts of Europe (Putaud et al., 2010, Schwartz et al., 2016). An assessment by Hendriks et al. (2013) found that secondary NH₄⁺ contributed 10 %–20 % of the PM_{2.5} mass in densely populated areas in Europe and even higher contributions in areas with intensive livestock farming. Concentrations of PM_{2.5} continue to exceed the EU limit values of 25 µg m⁻³ annual mean in large parts of Europe in 2017 (EEA, 2019). Particulate NH₄⁺ data presented from the DELTA[®] network therefore highlight the potential contribution of NH₃ of agricultural origin to fine NH₄⁺ aerosols in PM_{2.5}. The formation and transport of these secondary aerosols pose a serious risk to human health since PM_{2.5} is linked to increased mortality from respiratory and cardiopulmonary diseases (AQEG, 2012).

A considerable fraction of the aerosol components measured was made up of sea salt (Na⁺ and Cl⁻), with contributions from the sum of Na⁺ and Cl⁻ ranging from 4 % of the total aerosol loading at the inland Höglwald site in Germany (DE-Hog) to 43 % at Dripsey (IE-Dri), a coastal site in Ireland (Fig. 7). With the reduction in European emissions and concentrations of the gases SO₂, NO_x, and NH₃ for formation of NH₄⁺-containing aerosols, sea salt is therefore assuming a proportionate increase in the aerosol composition, consistent with observations from a recent European assessment of composition and trends in long-term EMEP measurements (EMEP, 2016). The concentrations of Ca²⁺ and Mg²⁺ were very low across the network, with values (mean of all sites < 0.1 µg m⁻³) that were at or below method limit of detection (LOD = ~ 0.1 µg m⁻³) (Table S3). These data are also considered to be underestimated due to the DELTA particle sampling cut-off (~ PM_{4.5}), and they were excluded from further assessment in this paper.

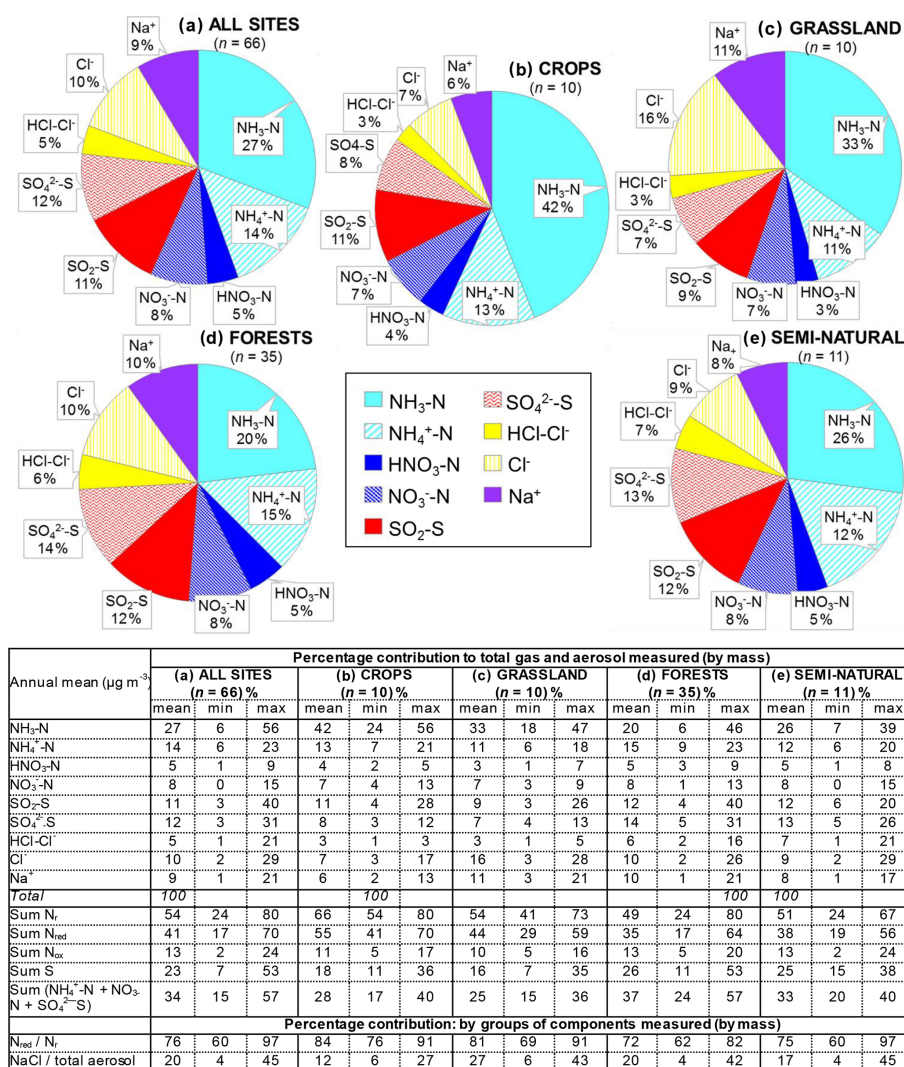


Figure 8. (a, b, c) Pie charts showing the mean atmospheric composition of gas and aerosol components from annual averaged concentrations ($\mu\text{g m}^{-3}$) measured at NEU DELTA[®] sites for (a) all sites ($n = 66$) and sites grouped according to ecosystem types, (b) crops ($n = 10$), (c) grassland ($n = 10$), (d) forests ($n = 35$), and (e) semi-natural ($n = 11$). UK-AMoP (parallel DELTA[®] at Auchencorth: NH₃ and NH₄⁺ only) and FR-FgsP (parallel DELTA[®] at Fougères: different sample train) were excluded in this analysis. (d, e) Summary statistics on percentage composition by mass ($\mu\text{g m}^{-3}$ element) measured. Sum N_r = sum (NH₃-N + NH₄⁺-N + HNO₃-N + NO₃⁻-N), sum S = sum (SO₂-S + SO₄²⁻-S), N_{red} = sum reduced N (NH₃-N + NH₄⁺-N), N_{ox} = sum oxidized N (HNO₃-N + NO₃⁻-N).

3.3.2 Comparisons with national gas emissions

In Fig. 9, the annual averaged gas and aerosol concentrations of grouped sites from each country are plotted with the corresponding national emission densities derived for NH₃, NO_x, and SO₂. The emissions data in the graphs are the 4-year averages for the period 2007 to 2010, expressed as emissions per unit area of the country per year ($\text{t km}^{-2} \text{ yr}^{-1}$) (see Sect. 2.6) and ranked in order of increasing emission densities. From the visual comparisons, national mean measured concentrations in each country appear to scale reasonably well with the ranked emission densities. This is supported by further regression analyses which showed significant cor-

relation between annual averaged concentrations of NH₃, NO_x, and SO₂ with emission densities of NH₃ ($R^2 = 0.49$, $p < 0.001$; Fig. 10a), NO_x ($R^2 = 0.20$, $p < 0.05$; Fig. 10b), and SO₂ ($R^2 = 0.65$, $p < 0.001$; Fig. 10c), respectively (Table 3). The particulate components NH₄⁺ and NO₃⁻ were also correlated with emission densities of NH₃ and HNO₃ (Table 3). By contrast, there was no relationship between SO₄²⁻ with emission densities of any of the three gases, possibly because of contributions to SO₄²⁻ from long-range transport. All regression plots of concentrations against emission densities, including summary statistics, are provided in Fig. S7.

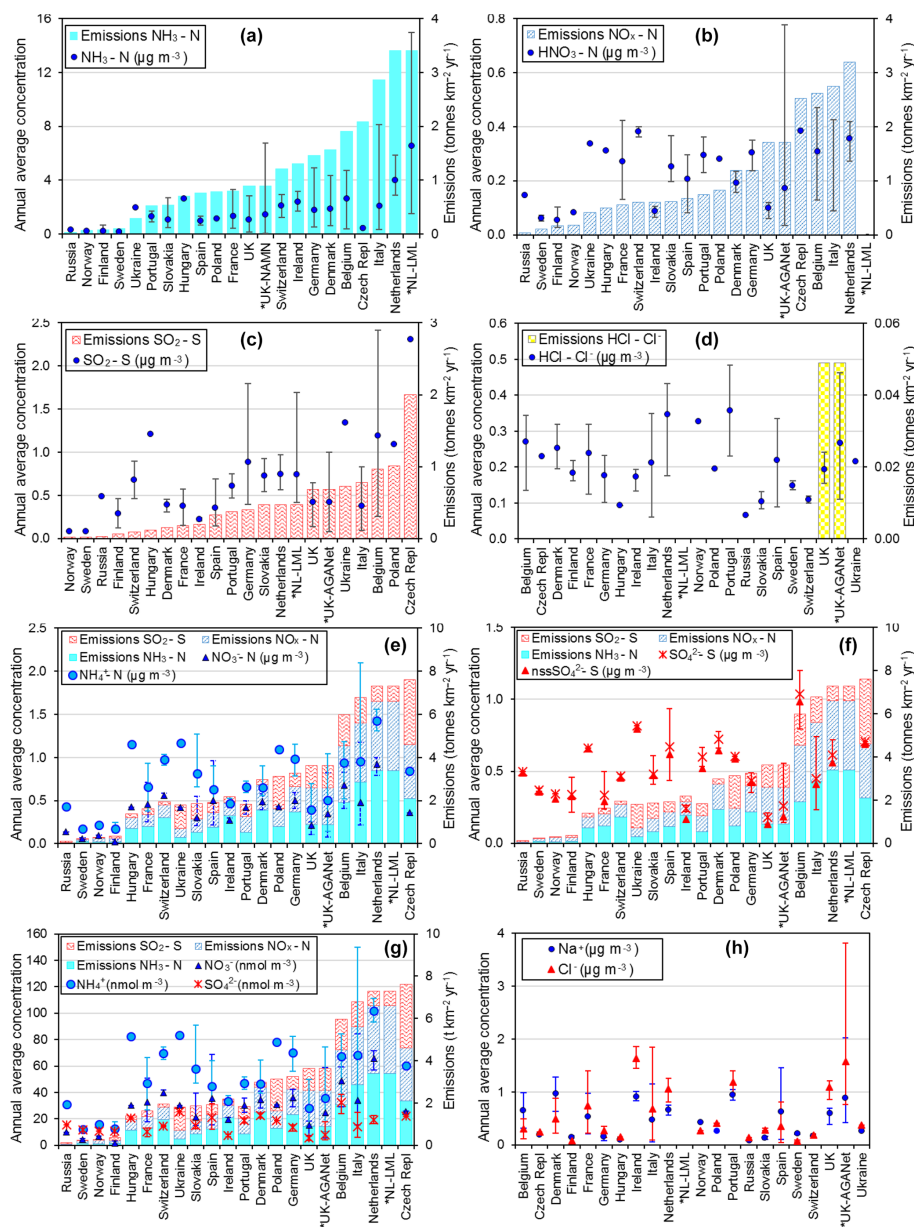


Figure 9. Comparisons of annual averaged gas and aerosol concentrations (2007–2010) of sites in the NEU DELTA[®] network, grouped by country, with the respective 4-year-averaged annual emission densities of gases (NH_3 , NO_x , and SO_2) over the same period. Monitoring data from three national monitoring networks: * UK NAMN (NH_3 from 72 sites and NH_4^+ from 30 sites; Tang et al., 2018a), * UK AGANet (raw uncorrected HNO_3 , SO_2 , HCl , NO_3^- , SO_4^{2-} , Cl^- , Na^+ from 30 sites; Tang et al., 2018b), and * NL-LML (NH_3 and SO_2 from 8 sites; van Zanten et al., 2017) are also included to illustrate the wider range of concentrations from larger numbers of sites. Error bars show the minimum and maximum concentrations measured in each country in the network. Where error bars are not visible, this indicates that either the country has measurement from just one site, or the range of concentrations measured are very close to the average.

The comparisons here used national emission totals, where emissions have been summed and averaged across very large and heterogeneous areas in each country. Additional analysis was also undertaken to compare the individual site mean data with (i) gridded emissions from individual $0.1^\circ \times 0.1^\circ$ EMEP grids in which the NEU sites are located (Figs. S8, S9) and (ii) averaged emissions of an extended number of

EMEP grids ($4 \times$ grids) closest to the site (Fig. S10). Since results from this analysis were similar to the comparisons with national emission densities, they are not included for further discussions in this paper. The purpose of the ranked emission densities is to compare the pollution climate in terms of primary gas emissions (SO_2 , NO_2 , NH_3) across the 20 European countries and to see if this is matched by the

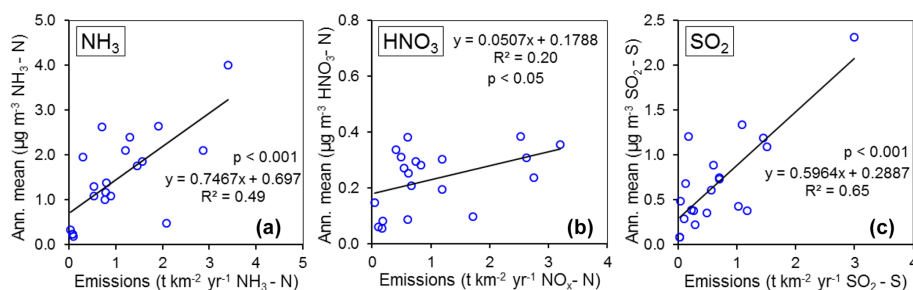


Figure 10. Regression plots of national annual averaged gas (NH₃, HNO₃, SO₂) concentrations (2007–2010) vs. 4-year national averaged emission densities of respective gases (NH₃, NO_x, and SO₂: t km⁻² yr⁻¹) from each country over the same period ($n = 20$).

Table 3. Summary statistics of regression analyses between national annual averaged gas (NH₃, HNO₃, SO₂) and aerosol (NH₄⁺, NO₃⁻, SO₄²⁻) concentrations and national emission densities (4-year average for period 2007 to 2010, expressed as emissions per unit area of the country per year) for each of the 20 countries in the NEU DELTA[®] network.

National annual average ($n = 20$) (µg m ⁻³)	National emission densities (20 countries)								
	NH ₃ (tonnes N km ⁻² yr ⁻¹)			NO _x (tonnes N km ⁻² yr ⁻¹)			SO ₂ (tonnes S km ⁻² yr ⁻¹)		
	Slope	Intercept	R ²	Slope	Intercept	R ²	Slope	Intercept	R ²
Gas NH ₃ -N	0.75	0.70	0.49***	0.57	0.90	0.30*	0.05	1.46	0.00 ^{ns}
Gas HNO ₃ -N	0.06	0.17	0.24*	0.05	0.18	0.20*	0.08	0.18	0.25*
Gas SO ₂ -S	0.17	0.52	0.24 ^{ns}	0.22	0.46	0.16 ^{ns}	0.60	0.29	0.65***
Aerosol NH ₄ -N	0.23	0.50	0.36**	0.19	0.54	0.27*	0.20	0.61	0.16 ^{ns}
Aerosol NO ₃ ⁻ -N	0.18	0.20	0.57***	0.15	0.23	0.44**	0.08	0.33	0.07 ^{ns}
Aerosol SO ₄ ²⁻ -S	0.06	0.47	0.07 ^{ns}	0.07	0.45	0.12 ^{ns}	0.12	0.44	0.18 ^{ns}

Significance level: * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$; ns: non-significant ($p > 0.05$).

DELTA[®] measurements. Despite the complex relationship between emissions and concentrations, the pollution gradient in Europe is clearly captured by the present data. At the same time, it also demonstrated the potential application of the DELTA[®] approach in providing national concentration fields as evidence to compare against spatial and long-term trends in the national emissions data.

3.3.3 Spatial variability across geographical regions

The form and concentrations of the different gas and aerosol components measured also varied according to geographic region across Europe (Fig. 11). The smallest concentrations (with the exception of SO₄²⁻ and Na⁺) were in northern Europe (Scandinavia), with broad elevations across other regions. Gas-phase NH₃ and particulate NH₄⁺ were the dominant species in all regions (Fig. 11). NH₃ showed the widest range of concentrations, with the largest concentrations in western Europe (mean = 2.4 NH₃ m⁻³, range = 0.2–7.1 µg NH₃ m⁻³, $n = 26$ in four countries). By contrast, HNO₃ and SO₂ concentrations were largest in high-NO_x- and high-SO₂-emitting countries in central and eastern Europe (Sect. 3.3.2). Particulate SO₄²⁻ concentrations were however more homogeneous between regions, which may

be attributed to atmospheric dispersion and long-range trans-boundary transport of this stable aerosol between countries in Europe (Szigeti et al., 2015; Schwarz et al., 2016). In the aerosol components, the spatial correlations between NO₃⁻, NH₄⁺, and NH₃ illustrate the potential for NH₃ emissions to drive the formation and thus regional variations in NH₄⁺ and NO₃⁻ aerosol. Particulate SO₄²⁻ concentrations in northern Europe (Scandinavia) were similar to other countries, despite having the smallest SO₂ and NH₃ emissions and concentrations (Fig. 9). By comparison, the smaller particulate NH₄⁺ and NO₃⁻ concentrations in northern Europe are consistent with the smallest emissions (NH₃ and NO_x) and concentrations of NH₃ and HNO₃ (Fig. 9). As discussed later in Sect. 3.4, the larger SO₄²⁻ concentrations reported in northern Europe were flagged up as anomalous from ion balance checks (ratio of NH₄⁺ : sum anions).

3.3.4 Comparisons by grouped components

In the following sections, variations in concentrations of the different gas and aerosol components according to ecosystem type (crops, grassland, forests, and semi-natural) and in relation to emissions (NH₃, NO_x, and SO₂) are further discussed. For ease of interpretation, components are grouped as

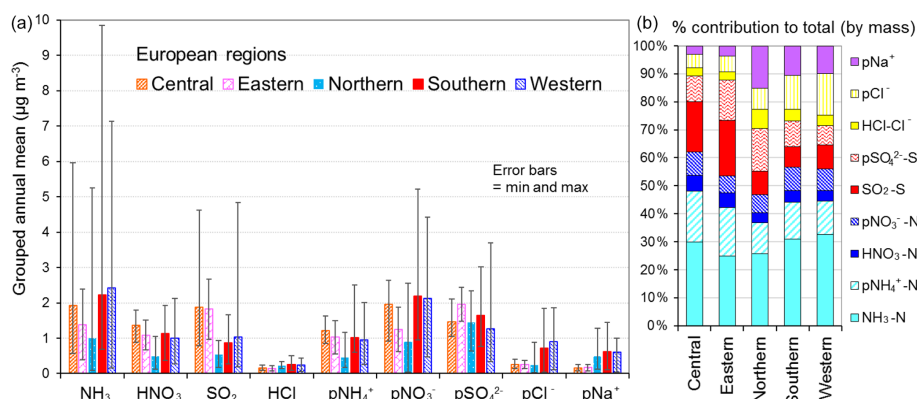


Figure 11. (a) Spatial variation in annual averaged gas and aerosol concentrations (2007 to 2010) measured in the NEU DELTA[®] network across Europe, grouped according to geographical distribution of the monitoring sites: central ($n = 17$), eastern ($n = 2$), northern ($n = 11$), southern ($n = 12$), and western ($n = 26$). The p in front of a component name denotes particulate. (b) Percentage composition of gas and aerosol components according to European region.

follows: reduced N (NH₃, NH₄⁺), oxidized N (HNO₃, NO₃[−]), S (SO₂, SO₄^{2−}), HCl, Na⁺, and Cl[−].

Reduced N (NH₃ and NH₄⁺)

Broad differences in NH₃ concentrations are observed between the grouped sites, with the largest concentrations at cropland sites, as expected, as these are intensively managed agricultural areas dominated by NH₃ emissions (Fig. 7a). Borgo Cioffi (IT-BCi) is an ecosystem station located in a 15 ha field (arable crops) on the Sele Plain, an agricultural area with intensive buffalo farming in southern Italy, and this provided the highest 4-year average of 8.1 µg NH₃–N m^{−3} (cf. group mean = 3.8 µg NH₃–N m^{−3}, $n = 10$) (Table 4, Table S9). Next highest in this group are the German Gebesee (DE-Geb) and the Belgian Lonzée (BE-Lon) sites, with 4-year-average concentrations of 4.9 and 4.8 µg NH₃–N m^{−3}, respectively (Table S9). At Gebesee, a decrease in NH₃ concentrations was observed over the 4-year period, falling almost 2-fold from an annual mean of 8.8 µg NH₃–N in 2007 to 4.8 µg NH₃–N in 2010 (Table S9). Annual mean concentrations in 2008 (2.9 µg NH₃–N m^{−3}) and 2009 (3.2 µg NH₃–N m^{−3}) were similar but smaller than in 2010. This illustrates the large inter-annual variability in concentrations that can occur even over a short time period. Variability between years may reflect the impact of changes in meteorological conditions on emissions from potential sources, with for example warmer, drier years increasing emissions and concentrations, contrasting with lower emissions and concentrations from the same source in a colder and wetter year. Episodic pollution events can also have a large influence on the annual mean concentration rather than the direct effects of changes in anthropogenic emissions over this short timescale. This suggests that for compliance assessment, an average over several years would provide a more robust basis than individual years. The assessment of trends also needs a longer

time series of at least 10 years (Tang et al., 2018a, b; Tørseth et al., 2012; van Zanten et al., 2017).

Grassland sites, with NH₃ emissions from grazing and fertilizers, provided the next highest concentrations, with annual averaged concentrations of 2.2 µg NH₃–N m^{−3} from the 10 sites in this group (Table 4). Cabauw in the Netherlands (NL-Cab) in this group was the second-highest NH₃ concentration site in the DELTA[®] network, after Borgo Cioffi (IT-BCi), with a 4-year-averaged annual concentration of 5.9 µg NH₃–N m^{−3} (Table S9). Unlike the Gebesee site (DE-Geb), annual NH₃ concentrations were consistent between years at Cabauw, ranging from annual mean of 6.3 µg NH₃–N m^{−3} in 2017 to 5.8 µg NH₃–N m^{−3} in 2010 (Table S9).

At the clean end of the NH₃ gradient are semi-natural and forest sites. The smallest concentrations were found at remote background sites in Russia (Fyodorovskoe bog, RU-Fyo) and the Scandinavian countries, in Finland (Lompola-jänkkä FI-Lom, Hyytiälä FI-Hyy, Sodankylä FI-Sod), Norway (Birkesnes, NO-Bir), and Sweden (Norunda SE-Nor, Skyytopr SE-Sky), where NH₃ concentration at each site was < 0.3 NH₃–N m^{−3} (Fig. 7, Table S9). By contrast, the semi-natural Horstermeer (NL-Hor) and forest sites Speulder (NL-Spe) and Loobos (NL-Loo) in the Netherlands gave concentrations that were 10-fold higher (2.9–4.1 µg NH₃–N m^{−3}) (Fig. 7, Table S9). This is consistent with much higher NH₃ emission density in the Netherlands (4-year average = 3.4 kt NH₃–N km^{−2} yr^{−1}) (Fig. 9).

With the exception of the Czech Republic, the annual averaged NH₃ concentrations scaled reasonably well with the 4-year-averaged mean NH₃ emission density in each country (Figs. 9, 10a1, 10b1) (see also Sect. 3.3.2). In the Czech Republic, measurement was made at a single site, BKFore (CZ-BK1), located at a remote forest location. The 4-year-averaged emissions in the EMEP grid (0.1° × 0.1°) containing the site are very small, at 2 t NH₃–N yr^{−1}, compared with an average of 68 t NH₃–N yr^{−1} (range = < 0.01

Table 4. Annual averaged concentrations of gas and aerosol concentrations, measured at all sites and at grouped sites classified according to each of the four ecosystem types in the NEU DELTA[®] network.

NEU Network	Annual averaged concentrations ($\mu\text{g m}^{-3}$) (2007–2010)								
	NH ₃ –N	NH ₄ –N	HNO ₃ –N	pNO ₃ [–] –N	SO ₂ –S	pSO ₄ ^{2–} –S	HCl–Cl [–]	Cl [–]	Na ⁺
All sites ($n = 66$)	1.63	0.73	0.23	0.42	0.58	0.48	0.22	0.57	0.46
Crops ($n = 10$)	3.81	1.11	0.32	0.61	0.87	0.63	0.24	0.58	0.49
Grassland ($n = 10$)	2.16	0.67	0.20	0.42	0.53	0.38	0.21	0.98	0.64
Forest ($n = 35$)	1.04	0.65	0.23	0.39	0.54	0.48	0.22	0.52	0.45
Semi-natural ($n = 11$)	1.11	0.70	0.18	0.35	0.50	0.43	0.22	0.37	0.30

to 567 t NH₃–N yr^{–1}) across the Czech Republic (Fig. S9). The low emissions, combined with the small concentrations measured at BKFore (0.5 $\mu\text{g NH}_3\text{–N m}^{-3}$), suggests it is highly likely to represent concentrations at the low end of the range of NH₃ concentrations that might be expected to be encountered in the Czech Republic. By comparison, Belgium has a similar emission density to the Czech Republic, but the mean concentrations from three sites (2.6 $\mu\text{g NH}_3\text{–N m}^{-3}$) encompassed sites located in cropland areas (Lonzée BE-Lon, 4.7 $\mu\text{g NH}_3\text{–N m}^{-3}$) and forest sites (Braschaat BE-Bra, 2.8 $\mu\text{g NH}_3\text{–N m}^{-3}$, and Vielsalm BE-Vie, 0.4 $\mu\text{g NH}_3\text{–N m}^{-3}$) (Table S9).

The markedly high concentrations of NH₃ across the NEU network indicates that contributions by emission and deposition of NH₃ would be a major contributor to the effects of N_r on sensitive habitats. In comparing the annual averaged NH₃ concentration with the revised UNECE “critical levels” of NH₃ concentrations (Cape et al., 2009), the lower limit of 1 $\mu\text{g NH}_3\text{ m}^{-3}$ annual mean for the protection of lichens and bryophytes was exceeded at 63 % of sites (40 sites in 15 countries) (Table S10). Even the higher 3 $\mu\text{g NH}_3\text{ m}^{-3}$ annual mean for the protection of vegetation was still exceeded at 27 % of sites (17 sites in 10 countries) (Table S10). Most notably, all four sites from the Netherlands were in exceedance of both the 1 and the 3 $\mu\text{g NH}_3\text{ m}^{-3}$ thresholds. The large concentrations in the Netherlands highlight the high levels of NH₃ that semi-natural and forest areas are exposed to within an intensive agricultural landscape, where 117 out of the 166 Natura 2000 areas were reported to be sensitive to nitrogen input (Lolkema et al., 2015). A recent assessment estimated that critical loads for eutrophication were exceeded in virtually all European countries and over about 62 % of the European ecosystem area in 2016 (EMEP, 2018). In particular, the highest exceedances occurred in the Po Valley (Italy), the Dutch–German–Danish border areas, and north-western Spain, where the highest NH₃ concentrations have been measured in this network. Since NH₃ is preferentially deposited to semi-natural and forests (high V_d to these ecosystem types; Sutton et al., 1995), then NH₃ will dominate dry N deposition and exert the larger ecological impact. In Flechard et al. (2011), dry NH₃–N deposition from the first 2 years of NH₃ measurement in the NEU DELTA[®] network was esti-

mated to contribute between 25 % and 50 % of total dry N deposition in forests, according to models. The fraction is larger in short semi-natural vegetation since V_d for NH₄⁺ and NO₃[–] is smaller in short vegetation than in forests (Flechard et al., 2011).

Comparison with NH₃ data from the Dutch LML network

The 4-year-averaged NH₃ concentrations from the Dutch LML air quality network (see Sect. 2.7.1) for the period 2007 to 2010 are plotted alongside the NH₃ measurements made at the four Dutch sites in the DELTA[®] network (Fig. 9a). The 4-year-averaged concentrations from the eight LML sites were between 1.5 and 15 $\mu\text{g NH}_3\text{–N m}^{-3}$, highlighting the high concentrations and spatial variability in concentrations in the Netherlands. The mean NH₃ concentrations measured at the four Dutch sites in the DELTA[®] network of 2.9 $\mu\text{g NH}_3\text{–N m}^{-3}$ (Horstermeer, NL-Hors; semi-natural) to 5.9 $\mu\text{g NH}_3\text{–N m}^{-3}$ (Cabauw, NL-Cab; grassland) were within the range of concentrations measured in the Dutch LML network.

Comparison with NH₃ data from the UK NAMN network

The 4-year-averaged NH₃ concentrations calculated from the 72 sites in the NAMN (see Sect. 2.7.2) for the period 2007 to 2010 were smaller than the Dutch LML network, ranging from 0.05 to 6.7 $\mu\text{g NH}_3\text{–N m}^{-3}$, consistent with smaller NH₃ emissions from the UK (Fig. 9a). In a joint collaboration between the UK and Dutch networks, inter-comparison of NH₃ measurements by the DELTA[®] method (monthly) with the AMOR wet chemistry system (hourly; van Zanten et al., 2017) was carried out at the Zegveld site (ID 633) in the Dutch LML network (van Zanten et al., 2017) between 2003 and 2015. Good agreement was provided, lending support for comparability between the independent measurements reported in Tang et al. (2018a).

Particulate NH₄⁺

Particulate NH₄⁺ concentrations across the 64 sites were more homogeneous than NH₃, varying over a narrower range

between $0.13 \mu\text{g NH}_4^+-\text{N m}^{-3}$ at Sodankylä (Finland, FI-Sod) and $2.1 \mu\text{g NH}_4^+-\text{N m}^{-3}$ at Borgo Cioffi (Italy, IT-BCi) (Fig. 7, Table S11). By comparison, the difference in NH₃ between the smallest ($0.07 \mu\text{g NH}_3-\text{N m}^{-3}$ at Lompolojänkää, Finland, FI-Lom) and largest ($8.1 \mu\text{g NH}_3-\text{N m}^{-3}$ at Borgo Cioffi, Italy, IT-BCi) concentrations varied by a factor of 110 (Fig. 7, Table S10). Secondary aerosols have longer atmospheric lifetimes and will therefore vary spatially much less than their precursor gas concentrations. While the concentrations of NH₃ vary at a local to regional level owing to large numbers of sources at ground level and high deposition in the landscape, NH₄⁺ is less influenced by proximity to NH₃ emission sources and varies in concentration at regional scales (Sutton et al., 1998; Tang et al., 2018a).

In Fig. 9, annual averaged NH₄⁺ concentrations ($\mu\text{g NH}_4^+-\text{N}$ in Fig. 9e; nmol m^{-3} in Fig. 9g) are plotted with 4-year-averaged emissions densities for NH₃, NO_x, and SO₂ from each country, with the combined total emission densities shown in ranked order. Regression analyses showed NH₄⁺ concentrations to be correlated with NH₃ emissions ($R^2 = 0.36$, $p < 0.01$, $n = 20$) and NO_x emissions ($R^2 = 0.27$, $p = 0.02$, $n = 20$) but not with SO₂ emissions (Table 3, Fig. S7). The smallest NH₄⁺ concentrations were in Sweden, Norway, and Finland (annual average $< 0.3 \mu\text{g NH}_4^+-\text{N m}^{-3}$), with the lowest emissions of NH₃, NO_x, and SO₂ and also the smallest concentrations of the precursors gases NH₃ ($< 0.3 \mu\text{g NH}_3-\text{N m}^{-3}$), HNO₃ ($< 0.1 \mu\text{g HNO}_3-\text{N m}^{-3}$), and SO₂ ($< 0.3 \mu\text{g SO}_2-\text{S m}^{-3}$).

The UK and Irish sites have the next smallest NH₄⁺ concentrations of 0.4 and $0.5 \mu\text{g NH}_4^+-\text{N m}^{-3}$ (cf. mean of all countries $= 0.74 \mu\text{g NH}_4^+-\text{N m}^{-3}$). Particulate NH₄⁺ data from the UK NAMN (Tang et al., 2018a) are also included for comparison. The 4-year-average concentrations from the 30 sites ($0.5 \mu\text{g NH}_4^+-\text{N m}^{-3}$, range $= 0.14$ to $1.0 \mu\text{g NH}_4^+-\text{N m}^{-3}$) are comparable with the mean of $0.40 \mu\text{g NH}_4^+-\text{N m}^{-3}$ (range $= 0.2$ to $0.9 \mu\text{g NH}_4^+-\text{N m}^{-3}$) from just 4 sites in the NEU network. A combination of lower emissions of precursor gases (Fig. 9) and being farther away from the influence of long-range transport of NH₄⁺ aerosols from the higher-emission countries on mainland Europe may be contributing factors to the small NH₄⁺ concentrations measured in the UK and Ireland.

The largest national mean concentration of particulate NH₄⁺ ($1.4 \mu\text{g NH}_4^+-\text{N m}^{-3}$) was measured in the Netherlands, which also has highest NH₃ and NO_x emissions (Fig. 9e). Indeed, the NH₄⁺ was matched by large NO₃⁻ concentration ($0.9 \mu\text{g HNO}_3-\text{N m}^{-3}$) (Fig. 9e), lending support to the contribution of NH₄NO₃ to the NH₄⁺ and NO₃⁻ load, together with contribution from (NH₄)₂SO₄ ($0.6 \mu\text{g SO}_4^{2-}-\text{S}$) (Fig. 9f). The particulate NH₄⁺ concentrations measured in Italy (mean $= 1.0 \mu\text{g NH}_4^+-\text{N m}^{-3}$) (Fig. 9e), which includes the site in the Po Valley (IT-PoV) with a mean concentration of $1.9 \mu\text{g NH}_4^+-\text{N m}^{-3}$ (Table S11), are comparable with

an assessment of PM_{2.5} composition at four sites in the Po Valley (Ricciardelli et al., 2017).

Oxidized N (HNO₃ and NO₃⁻)

The percentage mass contribution of oxidized N (sum of HNO₃ and NO₃⁻; $\mu\text{g N m}^{-3}$) to the total gas and aerosol species measured was on average 13 % (range $= 2\%$ – 24%) (Fig. 8). This compares with 41 % (range $= 17\%$ – 70%) from reduced N (sum NH₃ and NH₄⁺; $\mu\text{g N m}^{-3}$) and 23 % (range $= 7\%$ – 53%) from sulfur (sum of SO₂ and SO₄²⁻; $\mu\text{g S m}^{-3}$) (Fig. 8). DELTA[®] measurements of HNO₃ also include contributions from co-collected oxidized N species such as HONO (see Sect. 2.2.3) and are therefore an upper estimate that may in some cases be twice as large as the actual HNO₃ concentration, based on observations in the UK (Tang et al., 2018b; correction factor of 0.45) and from the parallel DELTA[®] measurements made at Fougères (FR-FgsP) (Fig. S5). At this site, HNO₃ measurement with NaCl-coated denuders provided an annual mean concentration of $0.08 \mu\text{g HNO}_3-\text{N m}^{-3}$, compared with $0.19 \mu\text{g HNO}_3-\text{N m}^{-3}$ measured on carbonate-coated denuders from the main site (FR-Fgs) (Table S7). With this caveat in mind, uncorrected annual mean HNO₃ concentrations were in the range of $0.03 \mu\text{g HNO}_3-\text{N}$ at Kaamenan (Finland; FI-Kaa) to $0.47 \mu\text{g HNO}_3-\text{N}$ at Braschaat (Belgium; BE-Bra) (Table S7). In Fig. 9b, HNO₃ concentrations are compared with NO_x emissions, the precursor gas for secondary formation of HNO₃. Overall, a weak but significant correlation was observed between concentrations of HNO₃ and NO_x emission densities across the 20 countries ($R^2 = 0.2$, $p < 0.05$) (Fig. 10b, Table 3, Fig. S2). Russia has the lowest NO_x emission densities ($0.04 \text{ t NO}_x-\text{N yr}^{-1}$), but HNO₃ from the single site ($0.15 \mu\text{g HNO}_3-\text{N m}^{-3}$) is larger than the smallest concentrations measured in Finland, Norway, and Sweden (annual average $< 0.1 \mu\text{g HNO}_3-\text{N m}^{-3}$). HNO₃ formation by photochemical processes may be enhanced in hotter, sunnier summer weather in Russia. Since SO₂ concentrations (mean $= 0.49 \mu\text{g SO}_2-\text{S}$) at the Russian site (RU-Fyo) are in molar excess over the low levels of NH₃ (mean $= 0.32 \mu\text{g NH}_3-\text{N m}^{-3}$), removal of HNO₃ by reaction with NH₃ will also be limited. HNO₃ concentrations in the UK and Ireland are marginally higher than the Scandinavian countries. Here, the annual averaged concentrations of HNO₃ are similar (0.10 vs. $0.09 \mu\text{g m}^{-3}$) (Table S12), despite NO_x emissions density ($\text{t km}^{-2} \text{ yr}^{-1}$) in the UK being 3 times larger than in Ireland (Fig. 9b). HNO₃ concentrations on the European continent were generally higher (0.2 – $0.4 \mu\text{g HNO}_3-\text{N m}^{-3}$).

In the UK, HNO₃ data are also available on a wider spatial scale from the AGANet (Sect. 2.7.2; Tang et al., 2018b). The 4-year-average concentrations of HNO₃ from 30 sites in the AGANet are plotted alongside the NEU HNO₃ data from the four UK sites in its network in Fig. 9b. The UK HNO₃ data on the UK-AIR database (<https://uk-air.defra.gov.uk/>, last ac-

cess: 25 November 2019) have been corrected for HONO interference with a 0.45 correction factor (see Tang et al., 2018b). For consistency in Fig. 9b, the UK raw uncorrected HNO₃ data are used for the present comparison. The 30-site mean ($0.17 \mu\text{g HNO}_3\text{-Nm}^{-3}$) was higher than from just 4 UK sites in the NEU network ($0.10 \mu\text{g HNO}_3\text{-Nm}^{-3}$). The range of concentrations was also wider, from $0.03 \mu\text{g HNO}_3\text{-Nm}^{-3}$ at a remote background site in Northern Ireland to $0.77 \mu\text{g HNO}_3\text{-Nm}^{-3}$ at a central London urban site, where interference from HONO and NO_x in HNO₃ determination is likely to be larger (Tang et al., 2015; 2018b).

Like particulate NH₄⁺, NO₃⁻ concentrations are also correlated with emission densities of NH₃ ($R^2 = 0.57$, $p < 0.001$, $n = 20$) and NO_x (slope = 0.15, $R^2 = 0.44$, $p < 0.01$, $n = 20$) but not with SO₂ (Table 3, Fig. S7). The smallest NO₃⁻ concentrations were again in Sweden, Norway, and Finland with low NH₃ and NO_x emissions and also smallest concentrations of HNO₃, SO₂, and NH₄⁺ in the network (Fig. 9). The largest NO₃⁻ concentrations were measured in the Netherlands, with a mean of $0.92 \mu\text{g NO}_3\text{-Nm}^{-3}$, compared with a network average of $0.39 \mu\text{g NO}_3\text{-Nm}^{-3}$ (Fig. 9e, Table S13). The higher NO₃⁻ concentrations correlated well with the high NH₃, HNO₃, and NH₄⁺ concentrations in the Netherlands (Fig. 9). This suggests that concentrations of NO₃⁻ are linked to local formation of NH₄NO₃, which is dependent on concentrations of NH₃ and HNO₃, and also to the influence of meteorology on transport of NH₄NO₃ between countries on mainland Europe and export out of Europe. Countries in Scandinavia such as Sweden, Norway, and Finland and in the British Isles are farthest from the influence of long-range transboundary transport from Europe, with concentrations of NH₄NO₃ that are smaller than on the continent.

Sulfur (SO₂ and SO₄²⁻)

Annual averaged SO₂ concentrations measured across the network were between 0.9 and $2.3 \mu\text{g SO}_2\text{-Sm}^{-3}$ (Fig. 9c, Table S14). By comparison, the EMEP network of 58 urban-background sites reported annual mean concentrations of 0.03 and $5.5 \mu\text{g SO}_2\text{-Sm}^{-3}$ over the same period, with the largest SO₂ concentrations from North Macedonia and Serbia. Since these high-emitting countries were not included in the DELTA[®] network, the range of SO₂ concentrations is smaller. Together, the small SO₂ concentrations reflect the substantial reductions in SO₂ emissions across Europe (−74 % between 1990 and 2010) and large reductions in ambient concentrations and deposition of sulfur species across Europe during the last decades (EMEP, 2016).

SO₂ concentrations were also correlated with SO₂ emission density ($R^2 = 0.65$, $p < 0.001$, $n = 20$) in each country (Fig. 10c, Table 3). The smallest and largest SO₂ annual average concentrations corresponded with the lowest emissions in Norway and highest in the Czech Republic (Fig. 9c). By contrast, SO₂ concentrations from the only measurement

site, Bugac, in Hungary (HU-Bug) are much higher than expected on the basis of SO₂ emission density estimated for the country. This suggests that Bugac is likely to be affected by proximity to sources. This contrasts with the BKFore site in the Czech Republic (CZ-BK1), which had smaller NH₃ concentrations due to its location away from sources.

Following emission, SO₂ disperses and undergoes chemical oxidation in the atmosphere to form SO₄²⁻ both in the gas phase and in cloud and rain droplets (Baek et al., 2004; Jones and Harrison, 2011). Particulate SO₄²⁻ produced is generally associated with NH₄⁺ and NO₃⁻ (see Sect. 3.4). The regional pattern of SO₄²⁻ was similar to, and correlated well with, particulate NH₄⁺ and NO₃⁻ (Fig. 9g), suggesting well-mixed air on the continent since (NH₄)₂SO₄ is stable and long-lived. Countries in the British Isles (UK and Ireland) and in Scandinavia (Sweden, Norway, Finland) have smaller concentrations of SO₄²⁻ (Table S15). They are located far enough away from sources and activities on continental Europe such that they are less influenced by the emissions from central Europe.

As discussed earlier, particulate NH₄⁺ and NO₃⁻ concentrations were smallest in the Scandinavian countries, which corresponded with low emission densities of the precursor gases NH₃ and NO_x. By analogy, since these countries also have the lowest emission densities of SO₂ (Fig. 9c), particulate SO₄²⁻ concentrations would be expected to be similarly low. Particulate SO₄²⁻ in Finland and Norway (mean = $0.34 \mu\text{g SO}_4\text{-Sm}^{-3}$) and Sweden (mean = $0.37 \mu\text{g SO}_4\text{-Sm}^{-3}$) were however comparable with concentrations on mainland Europe (range = 0.33 to $1.0 \mu\text{g SO}_4\text{-Sm}^{-3}$) and larger than the UK ($0.18 \mu\text{g SO}_4\text{-Sm}^{-3}$) and Ireland ($0.24 \mu\text{g SO}_4\text{-Sm}^{-3}$) (Fig. 9f). An ion balance check on the ratio of equivalent concentrations of NH₄⁺ to the sum of NO₃⁻ and SO₄²⁻ (see Sect. 3.4) was less than 0.5. Since NH₄⁺ is a counter-ion to NO₃⁻ and SO₄²⁻ formation, the imbalance suggests that SO₄²⁻ concentrations may be overestimated at the sites in Sweden, Norway, and Finland.

HCl, Cl⁻, and Na⁺

The average concentrations of HCl across the network were of low magnitude, with limited variability, ranging from 0.07 in Russia to $0.36 \mu\text{g HCl-Cl}^{-}\text{m}^{-3}$ in Portugal (Fig. 9d). At a site level, HCl concentrations varied between 0.06 at Renon (Italy; IT-Ren – inland location) to $0.48 \mu\text{g HCl-Cl}^{-}\text{m}^{-3}$ at Espirra (Portugal, PT-Esp – coastal location) (Table S16). In the UK AGANet network, the highest concentrations of HCl were found in the source areas in the south-east and south-west of England and also in central England, north of a large coal-fired power station (Tang et al., 2018b). HCl emissions and concentrations in the atmosphere are mostly derived from combustion of fossil fuels (coal and oil), biomass burning, and the burning of municipal and domestic waste

in municipal incinerators (Roth and Okada 1998; McCulloch et al., 1999; Ianniello et al., 2011). Several manufacturing processes, including cement production, also emit HCl (McCulloch et al., 1999). At coastal sites, HCl released from the reaction of sea salt with HNO₃ and H₂SO₄ can be a significant source (Roth and Okada 1998; Keene et al., 1999; McCulloch et al., 1999; Ianniello et al., 2011). UK is the only country with available HCl emission estimates (<https://naei.beis.gov.uk/data/>, last access: 3 January 2020). Emissions of HCl in the UK (mainly from coal burning in power stations) have declined to very low levels, from 74 kt in 1999 to 5.7 kt in 2015. The 4-year-averaged emission density for HCl for the period 2007 to 2010 was just 0.05 tonnes HCl–Cl[–] km^{–2} yr^{–1}, although HCl emissions could still pose a threat to sensitive habitats close to sources (Evans et al., 2011). The low HCl concentrations measured in the network would suggest that the shift in Europe's energy system from coal to other sources has contributed to low HCl emissions (UK) and concentrations (observed across the network).

Particulate Cl[–] on the other hand is predominantly marine in origin, with sea salt (NaCl) as the most significant source (Keene et al., 1999). Molar concentrations of Cl[–] and Na⁺ are seen to be similar in most countries, demonstrating close coupling between the two components (Fig. 9h). The largest concentrations of Na⁺ and Cl[–] occurred in coastal countries such as the UK, Ireland, Netherlands, and Portugal, with the highest of country-averaged annual concentrations of 1.6 µg Cl[–] m^{–3} and 0.9 µg Na⁺ m^{–3} from Ireland (Tables S16 and S17). Data from the 30 sites in the UK AGANet network showed a wider range of Cl[–] and Na⁺ concentrations (Fig. 9h). The AGANet site with the largest 4-year-averaged annual concentrations of 3.8 µg Cl m^{–3} and 2.0 µg Na⁺ m^{–3} is Lerwick from the east coast of the Shetland Islands, exposed to the North Atlantic.

Farther away from the coastal influence of marine aerosol, the smallest concentrations of Cl[–] and Na⁺ were measured in landlocked countries such as Germany (mean of all sites = 0.27 µg Cl[–] m^{–3} and 0.15 µg Na⁺ m^{–3}). Concentrations in Hungary, Poland, the Czech Republic, and Russia were also low, but inferences about these countries are necessarily limited by measurements at a single site in each of these countries. At coastal sites in Norway (NO-Bir) and Sweden (SE-Nor and SE-Sk2), the very low particulate Cl[–] concentrations (< 0.1–0.3 µg m^{–3}) and high Na : Cl molar ratios (3–5) are anomalous. It is possible for sea salt to be depleted in Cl[–] (through the loss of HCl gas) by the reaction of NaCl particles with atmospheric acids (Finlayson-Pitts and Pitts, 1999; Keene et al., 1999), leading to high Na : Cl ratios for sea salts transported over long distances. The coastal locations of these sites (Fig. 2) suggest that they are more likely to be influenced by freshly generated marine aerosols (cf. coastal sites in UK and Ireland), and larger concentrations of sea salt (Na⁺ and Cl[–]) and a 1 : 1 relationship between Na⁺ and Cl[–] are expected. The Cl[–] concentrations

are likely to be underestimated at these sites (see Sect. 3.2.3) and are further discussed in the next section (Sect. 3.4).

3.4 Correlations between gas and aerosol components

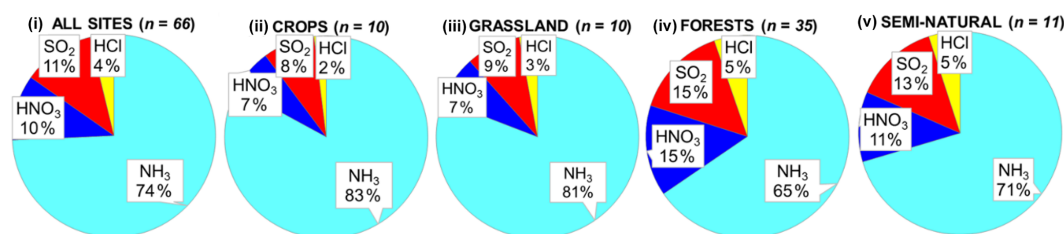
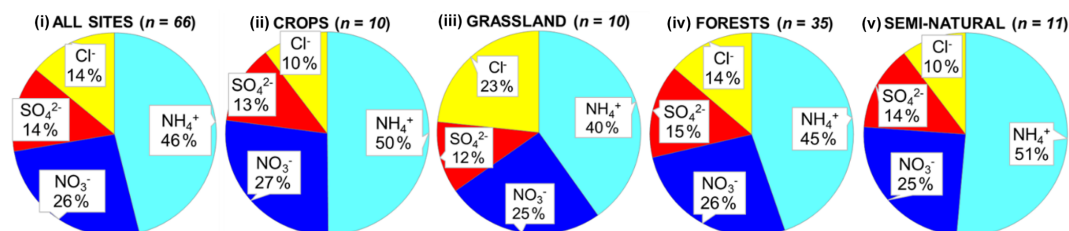
Regression analyses were carried out between the mean molar-equivalent concentrations of all inorganic gas and aerosol components measured at each site ($n = 66$; Fr-FgsP and UK-AMoP excluded) in the NEU network, with summary statistics provided in Table 5. With the exception of SO₂ vs. HCl ($R^2 = 0.05$, $p > 0.05$), the gases were positively correlated with each other, possibly due to similarities in the regional distribution of their emissions and concentrations. Comparing the mean molar concentrations of NH₃ with SO₂ and HNO₃ showed that NH₃ was on average 6-fold and 7-fold higher, respectively, whereas molar concentrations of SO₂ and HNO₃ were similar (Table 6, Fig. 12). The molar ratio of NH₃ to the sum of all acid gases (SO₂, HNO₃, and HCl) was on average 3 (Table 6, Fig. 12), confirming that there is a surplus of the alkaline NH₃ gas to neutralize the atmospheric acids in the atmosphere, similar to that observed in the UK (Tang et al., 2018b). With the more substantial decline in emissions of SO₂, compared with a more modest reduction in NO_x, the concentrations of SO₂ are at a level where it is no longer the dominant acid gas, such that HNO₃ and HCl are together contributing a larger fraction of the total acidity in the atmosphere in the present assessment.

In the aerosol phase, NH₄⁺ correlated well with NO₃[–] ($R^2 = 0.75$, $p < 0.001$; Fig. 13a) and SO₄^{2–} ($R^2 = 0.75$, $p < 0.001$; Fig. 13b) (Tables 5 and 7) but not with Cl[–] (Table 5). Regression of the molar-equivalent concentrations of the sum of NO₃[–] and SO₄^{2–} against NH₄⁺ shows points close to the 1 : 1 line (slope = 0.84) and significant correlation ($R^2 = 0.64$, $p < 0.001$), which demonstrates the close coupling between the base NH₄⁺ and the acid NO₃[–] + SO₄^{2–} aerosols (Fig. 13c, Table 7). The reaction of NH₃ with H₂SO₄ is irreversible (i.e. “one-way”) under atmospheric conditions (Baek et al., 2004; Finlayson-Pitts and Pitts, 1999; Jones and Harrison, 2011; Huntzicker et al., 1980), whereas any NH₄NO₃ or NH₄Cl that is formed can dissociate to release NH₃, which can then be “removed” by reaction with H₂SO₄. The lack of correlation between NH₄⁺ and Cl[–] ($R^2 = 0.00$; Table 5) in the analysis suggests that NH₄⁺ is mainly associated with NO₃[–] and SO₄^{2–}.

Particulate Cl[–] was correlated with Na⁺ ($R^2 = 0.65$, $p < 0.001$) (Fig. 13f, Tables 5, 7), consistent with observations that NaCl in atmospheric aerosols is mainly sea salt in origin (O'Dowd and de Leeuw, 2007; Tang et al., 2018b). Like the precursor gases, the molar concentrations of particulate NH₄⁺ are larger than either NO₃[–] or SO₄^{2–} (Fig. 12, Table 8). Particulate NO₃[–] concentrations were on average 2-fold higher than particulate SO₄^{2–} (on a molar basis) so that there was twice as much NH₄NO₃ (Fig. 13a) as (NH₄)₂SO₄ (Fig. 13b). The shift in PM composition from (NH₄)₂SO₄ to NH₄NO₃ across Europe is well documented (Bleeker et al.,

Table 5. Regression correlations (R^2) between the mean molar concentrations (nmol m^{-3}) of gas and aerosol components at sites ($n = 66$) in the NEU DELTA[®] network.

	HNO ₃	HCl	SO ₂	NH ₃	NO ₃ ⁻	Cl ⁻	2 × SO ₄ ²⁻	2 × nss-SO ₄ ²⁻	NH ₄ ⁺	Na ⁺
HNO ₃	1									
HCl	0.13**	1								
SO ₂	0.46***	0.05 ^{ns}	1							
NH ₃	0.28***	0.11**	0.08*	1						
NO ₃ ⁻	0.66***	0.21**	0.19***	0.43***	1					
Cl ⁻	0.00 ^{ns}	0.22***	0.01 ^{ns}	0.11**	0.06*	1				
2 × SO ₄ ²⁻	0.34***	0.24***	0.33***	0.18***	0.39***	0.01 ^{ns}	1			
2 × nss-SO ₄ ²⁻	0.35***	0.17***	0.36***	0.15**	0.35***	0.04 ^{ns}	0.98***	1		
NH ₄ ⁺	0.72***	0.06 ^{ns}	0.34***	0.43***	0.75***	0.00 ^{ns}	0.28***	0.30***	1	
Na ⁺	0.00 ^{ns}	0.42***	0.00 ^{ns}	0.10**	0.13**	0.65***	0.09*	0.03 ^{ns}	0.00 ^{ns}	1

Significance level: * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$; ns: non-significant ($p > 0.05$).**(a) Gases: % contribution to total (sum of NH₃, HNO₃, SO₂, HCl) (nmol m^{-3})****(b) Aerosols: % contribution to total (sum of NH₄⁺, NO₃⁻, SO₄²⁻, Cl⁻) (nmol m^{-3})****Figure 12.** Pie charts of mean relative proportions of (a) gases (NH₃, HNO₃, SO₂, HCl) and (b) aerosols (NH₄⁺, NO₃⁻, SO₄²⁻, Cl⁻). Data are annual averaged concentrations (nmol m^{-3}) measured at NEU DELTA[®] sites for (i) all sites ($n = 66$) and sites grouped according to ecosystem types, (ii) crops ($n = 10$), (iii) grassland ($n = 10$), (iv) forests ($n = 35$), and (v) semi-natural ($n = 11$). UK-AMoP (parallel DELTA[®] at Auchencorth: NH₃ and NH₄⁺ only) and FR-FgsP (parallel DELTA[®] at Fougères: different sample train) were excluded in this analysis.

2009; Fowler et al., 2009; Tang et al., 2018b; Tørseth et al., 2012).

Non-sea-salt SO₄²⁻ (nss-SO₄²⁻) was also estimated from the SO₄²⁻ and Na⁺ data (see Sect. 2.2.1). The nss-SO₄²⁻ is estimated to comprise on average 25 % (range = 3 %–83 %, $n = 187$) of the measured total SO₄²⁻ aerosol (Table 8). This demonstrates that sea salt SO₄²⁻ (ss-SO₄²⁻) aerosol makes up a large and variable fraction of the total SO₄²⁻ measured, consistent with observations of the contribution by ss-SO₄²⁻ to the total SO₄²⁻ in precipitation observed in the wet-deposition measurements in this study (Fig. 11) and across Europe (ROTAP, 2012). Regression of nss-SO₄²⁻ vs.

NH₄⁺ (slope = 0.27, $R^2 = 0.30$) was not significantly different from the regression of SO₄²⁻ vs. NH₄⁺ (slope = 0.27, $R^2 = 0.28$) (Table 5). This suggests that NH₄⁺ is mainly associated with the nss-SO₄²⁻.

Correlation between NH₄⁺ and the sum of anions (NO₃⁻ + SO₄²⁻) is an important point of discussion (Table 7) as the ion balance serves as a quality check for the aerosol measurement. Due to some outliers in the comparison, the correlation between NH₄⁺ and SO₄²⁻ ($R^2 = 0.28$; Fig. 13b) is weaker than between NH₄⁺ and NO₃⁻ ($R^2 = 0.75$; Fig. 13c, Table 7). The outliers were measurements made by NILU and CEAM, although these vary according to monitor-

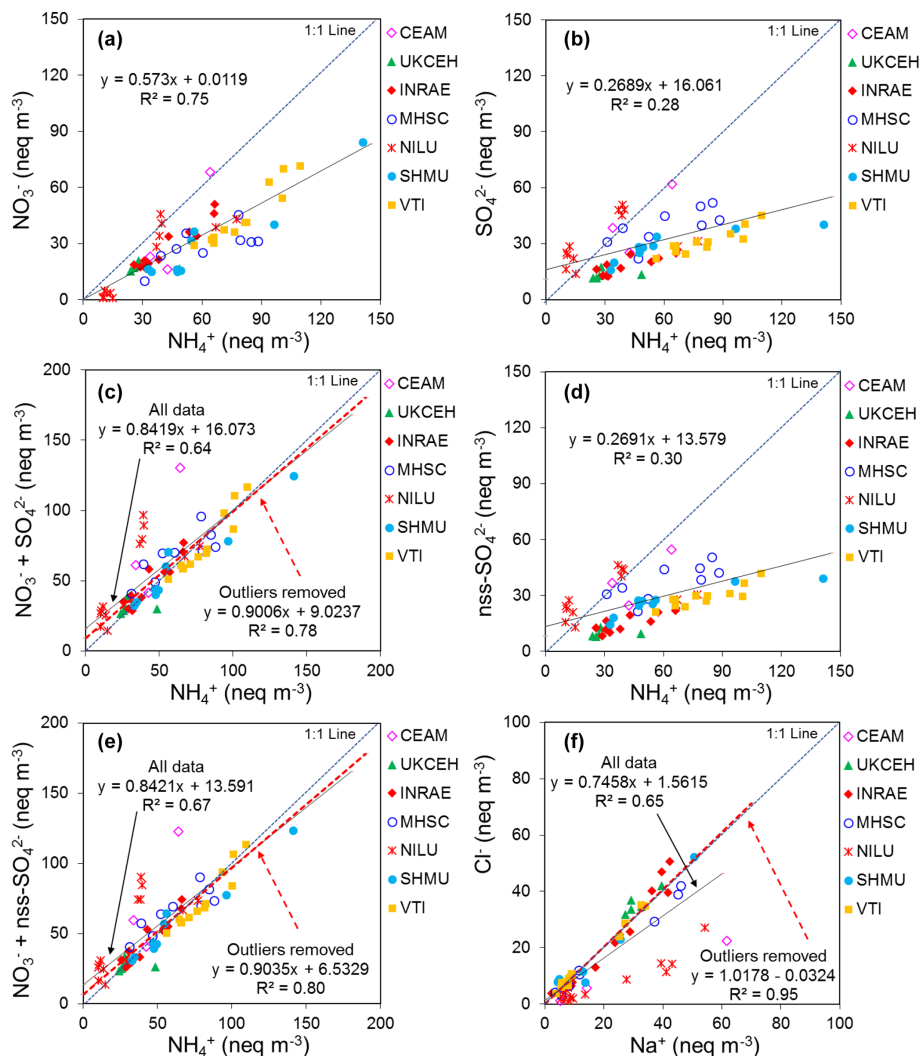


Figure 13. Regression plots between mean molar-equivalent concentrations of (a) NH_4^+ and NO_3^- , (b) NH_4^+ and SO_4^{2-} , (c) NH_4^+ and sum ($\text{NO}_3^- + \text{SO}_4^{2-}$), (d) NH_4^+ and nss-SO_4^{2-} , (e) NH_4^+ and sum ($\text{NO}_3^- + \text{nss-SO}_4^{2-}$), and (f) Na^+ and Cl^- , measured in the NEU DELTA[®] network. Each data point represents the mean of all monthly measurements at each site, with different coloured symbols for each laboratory making the measurements. Outliers: where equivalent concentrations of NH_4^+ : sum (anions) < 0.5 and $\text{Na}^+ : \text{Cl}^- > 2$.

Table 6. Mean molar concentrations of gases and NH_3 : acid gas ratios measured at sites ($n = 66$) in the NEU DELTA[®] network.

All NEU sites	Molar concentrations (nmol m^{-3})					Ratios		
	NH_3	HNO_3	SO_2	HCl	Sum acids	$\text{NH}_3 : \text{HNO}_3$	$\text{NH}_3 : \text{SO}_2$	$\text{NH}_3 : \text{sum acids}$
Mean	115	16.5	18.3	6.4	41.1	7.5	7.7	2.9
Min	5.4	2.0	2.5	1.6	10.9	0.8	0.5	0.3
Max	566	33.8	78.2	13.4	122	34	33	13
SD	108	8.4	14.7	2.8	22.4	7.2	6.6	2.6
n	66	66	66	66	66	66	66	66

Table 7. Linear regressions between the mean molar-equivalent concentrations of aerosol components (nanoequivalent, neq m⁻³) at sites (*n* = 66) in the NEU DELTA[®] network.

Linear regression	Mean molar-equivalent concentrations (neq m ⁻³)						
	NH ₄ ⁺ vs. NO ₃ ⁻	NH ₄ ⁺ vs. SO ₄ ²⁻	NH ₄ ⁺ vs. sum (NO ₃ ⁻ + SO ₄ ²⁻)	Na ⁺ vs. nss-SO ₄ ²⁻	NH ₄ ⁺ vs. sum (NO ₃ ⁻ + nss-SO ₄ ²⁻)	Na ⁺ vs. Cl ⁻ (all data)	Na ⁺ vs. Cl ⁻ (outliers excluded)
<i>R</i> ²	0.75***	0.28***	0.64***	0.30***	0.67***	0.65***	0.95***
Slope	0.57***	0.27***	0.84 ^{ns}	0.27***	0.84*	0.75***	1.01 ^{ns}
Intercept	0.01 ^{ns}	16.1***	16.1**	13.6***	13.6**	1.56 ^{ns}	-0.05 ^{ns}
No. of sites: <i>n</i>	66	66	66	66	66	66	50

Significance level: * *p* < 0.05, ** *p* < 0.01, *** *p* < 0.001; ns: non-significant (*p* > 0.05).

Table 8. Mean molar concentrations of aerosols and ratios measured at sites (*n* = 66) in the NEU DELTA[®] network.

All NEU sites	Molar concentrations (nmol m ⁻³)				Ratios			
	NH ₄ ⁺	NO ₃ ⁻	SO ₄ ²⁻	nss-SO ₄ ²⁻	NH ₄ ⁺ : NO ₃ ⁻	NH ₄ ⁺ : 2 × SO ₄ ²⁻	NH ₄ ⁺ : 2 × nss-SO ₄ ²⁻	NH ₄ ⁺ : (NO ₃ ⁻ + 2 × SO ₄ ²⁻)
Mean	52.8	30.2	15.1	13.9	2.4	1.8	2.1	0.9
Min	10.1	0.7	5.8	4	0.9	0.4	0.4	0.4
Max	141	84.3	38.4	35.8	21	3.6	5.1	1.6
SD	27.6	18.2	7.0	6.8	2.7	0.8	0.9	0.3
<i>n</i>	66	66	66	66	66	66	66	66

ing locations. The NILU laboratory made DELTA[®] measurements for 16 sites in six different countries (Belgium, Denmark, Finland, Norway, Sweden, and Switzerland). At three sites (Kaamanen, FI-Kaa; Laegern, CH-Lae; Oensingen, CH-Oe1), the ion balance of equivalent concentrations of NH₄⁺ : sum (NO₃⁻ + SO₄²⁻) was 1.0, whereas the ratios at the other 13 sites were between 0.4 and 0.7. The CEAM laboratory made measurements for all three sites in Spain. For CEAM, the ion balance ratio at Vall de Aliñá (ES-VDA) was 1, whereas the other two sites had ratios of 0.5 and 0.6.

Removal of the outlier NILU (7 out of 16) and CEAM (1 out of 3) data points with ion balance ratio < 0.5 improved both the slope (new slope = 0.90) and correlation (new *R*² = 0.78) (Fig. 13c). This indicates either an over-read of the anions (NO₃⁻, SO₄²⁻) or under-read of NH₄⁺ concentrations by the two laboratories at some sites. Results reported by NILU in the DELTA[®] field inter-comparisons (Sect. 3.2) showed that, with the exception of a few high NH₄⁺ and NO₃⁻ readings, there was on average no overall bias in the NH₄⁺, NO₃⁻, or SO₄²⁻ measurements by the NILU laboratory that could account for the high SO₄²⁻ outliers in the regression (Fig. 13). An inspection of individual monthly site data reported by NILU showed that 15 % of aerosol NH₄⁺ and 17 % of NO₃⁻ concentrations were below 0.1 µg m⁻³, compared with only 0.7 % of all SO₄²⁻ data. This then points to a potential under-read in NH₄⁺ and NO₃⁻. Possible reasons include

- loss of NH₄⁺, NO₃⁻ from filters (e.g. microbial degradation);
- non-capture on the aerosol filters (e.g. aerosol filters installed wrong way around);

iii. filters mixed up and wrong analysis performed on the acid- and base-coated filters;

iv. high blanks subtracted from already low concentrations at clean sites.

Possibilities still remain, however, of a potential over-read of SO₄²⁻. The ion balance checks suggest increased uncertainty in the NH₄⁺, NO₃⁻, and SO₄²⁻ measurements for seven sites: Hyytiälä (FI-Hyy), Sodankylä (FI-Sod), Rimi (DK-Rim), Risbyholm (DK-Ris), Soroe (DK-Sor), Skytöpp (SE-Sk2), and Vielsalm (BE-Vie). Examination of monthly site data from CEAM showed only 1.5 % of aerosol NH₄⁺ and 0.8 % of NO₃⁻ concentrations below 0.1 µg m⁻³, whereas all SO₄²⁻ data were above 0.1 µg m⁻³. For the CEAM lab, the uncertainty in NH₄⁺, NO₃⁻, and SO₄²⁻ measurements affected two sites: El Saler (ES-El) and Las Majadas (ES-Lam) (see also Sect. 3.3.3).

The regression of Na⁺ and Cl⁻ also showed the majority of data points close to the 1 : 1 line but with a small group of outliers below the 1 : 1 line from the CEAM and NILU laboratories (Fig. 13f). Both laboratories performed well in laboratory PT schemes (Sect. 3.1), with more than 80 % of reported data agreeing within ±10 % of reference values in both Na⁺ and Cl⁻ and no bias in the analytical method. The outliers in the ion balance therefore suggests some problems with Na⁺ and Cl⁻ determination on the DELTA[®] aerosol filters. Na⁺ and Cl⁻ data for some of the field DELTA[®] inter-comparisons were omitted from submissions by CEAM and NILU, and submitted data were in poor agreement with other laboratories (Sect. 3.2). Further regression analyses were carried out on individual monthly data, with sites grouped ac-

cording to measurements made by each of the seven laboratories (Fig. S11). Regressions for CEAM and NILU show the vast majority of data points below the 1 : 1 line, indicating a systematic underestimation of particulate Cl^- concentrations. The other five laboratories (INRAE, MHSC, SHMU, UKCEH, and vTI) all have data points close to the 1 : 1 line, with larger scatter both above and below the 1 : 1 line at lower concentrations. In Fig. 13f, a new regression line has therefore also been fitted, where outlier data with Na : Cl ratios > 2 from NILU (13 out of 16 sites) and CEAM (all 3 sites) have been removed. Exclusion of the outlier data points provided a regression line that is not significantly different from unity (slope = 1.02), with an R^2 value of 0.95 ($p < 0.001$). The near-1 : 1 relationship between particulate Na^+ and Cl^- is consistent with their origin from sea salt (NaCl).

The ion balance checks, together with the regular PT exercises and field inter-comparisons, therefore provided the platform against which to assess data quality and comparability of measurements between laboratories. This shows that overall, with the exception of a few identified outlier measurements, the laboratories are performing well and providing good agreement.

3.5 Seasonal variability in gases and aerosol

The time series of monthly averaged concentrations for the period 2006 to 2010 have been plotted to examine seasonality in the different gas and aerosol components according to ecosystem types (crops, grassland, semi-natural, and forests) (Fig. 14) and geographical regions (Fig. 15). Distinct seasonality was observed in the data, influenced by seasonal changes in emissions, chemical interactions, and the influence of meteorology on partitioning between the main inorganic gases and aerosol species.

3.5.1 NH_3

Distinctive and contrasting features in the seasonal cycle are observed, with the largest concentrations at cropland sites and smallest at semi-natural and forest sites (Fig. 14a). Similar to those observed in the annual mean concentrations (Fig. 9, 11), the monthly concentrations are also smallest in northern Europe and largest in western Europe (Fig. 15a).

Semi-natural sites

There are two distinct peaks in the seasonal cycle of grouped semi-natural sites: in April (mean = $2.2 \mu\text{g NH}_3 \text{ m}^{-3}$, $n = 12$) and in July (mean = $1.9 \mu\text{g NH}_3 \text{ m}^{-3}$, $n = 12$) (Fig. 14a). Since these sites are located away from agricultural sources, the seasonality in NH_3 concentrations is mostly governed by changes in environmental conditions and regional changes in NH_3 emissions. The differences in concentrations between the summer and winter at these sites were by a factor of 3, with the smallest concentrations in wintertime (December

and January), when low temperatures and wetter conditions decrease NH_3 emissions from regional agricultural sources while favouring a thermodynamic shift from gaseous NH_3 to the aerosol NH_4^+ phase. Conversely, warm, dry conditions in summer increases surface volatilization of NH_3 from low-density grazing livestock and wild animals and favour a thermodynamic shift to the gaseous (NH_3) phase, producing the summer peak. Vegetation is another potential source at these background sites under the right conditions (Flecharde et al., 2013; Massad et al., 2010). A complex interaction between atmospheric NH_3 concentrations and vegetation can lead to both emission and deposition fluxes, known as “bidirectional exchange”, dependent on relative differences in concentrations. This process is controlled by the so-called “compensation point”, defined as the concentration below which growing plants start to emit NH_3 into the atmosphere (Flecharde et al., 1999; Massad et al., 2010; Sutton et al., 1995). At sites distant from intensive farming and emissions, the bidirectional exchange with vegetation will partly control NH_3 concentrations. Inclusion of bidirectional exchange in dispersion modelling of NH_3 by incorporating a “canopy compensation point” is shown to improve model results for NH_3 concentrations in remote areas (e.g. Smith et al., 2000; Flecharde et al., 1999, 2011; Massad et al., 2010). The larger peak in April at these sites on the other hand suggests the influence of emissions from agricultural sources, e.g. from land spreading of manures.

Forest sites

The average seasonal cycle from the forest sites is similar to that of the semi-natural sites but diverged over the summer months (Fig. 14a). Here, the seasonal profile is characterized by the absence of any peaks in summer, with concentrations plateauing between May and August. Studies have shown that atmospherically deposited N is taken up by forest canopies since growth in forest ecosystems is commonly limited by the availability of N (Sievering et al., 2007), and tree canopies are a potential sink for atmospheric NH_3 (Fowler et al., 1989; Theobald et al., 2001). The capture and uptake of NH_3 during the growing seasons over the summer period could therefore account for the absence of a summer peak in NH_3 concentrations at forest monitoring sites, although a similar effect would also be expected for semi-natural sites.

Cropland sites

Fertilizers and arable crops are significant sources of NH_3 emissions and concentrations in an intensive agricultural landscape. Sites in this group showed considerably higher monthly mean monitored NH_3 concentrations than the other groups (Fig. 14a). A more complex seasonal pattern can be seen, with three peaks in NH_3 concentrations. Concentrations here are also lowest in the winter, although the wintertime concentrations are 3 times larger than semi-natural and

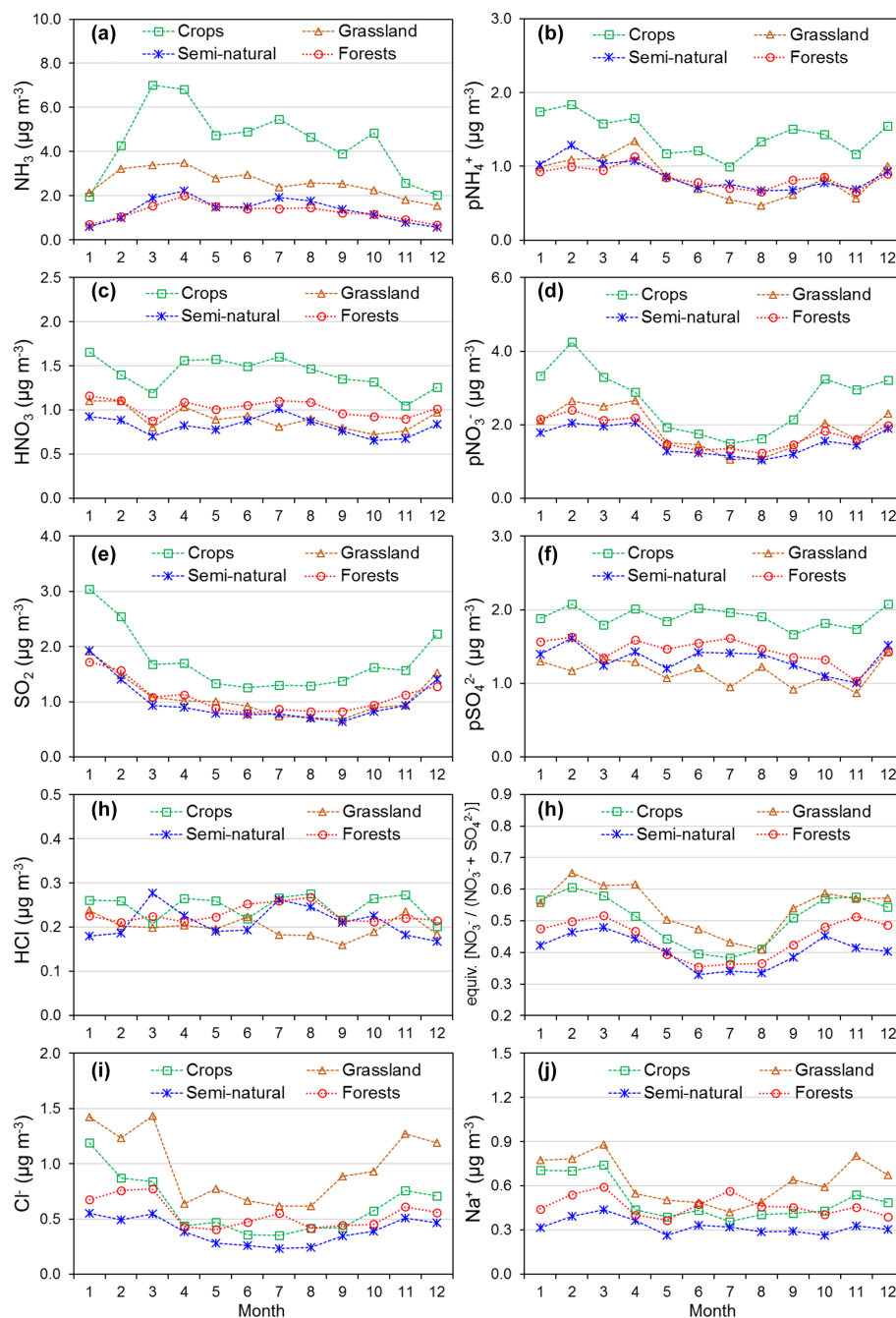


Figure 14. Seasonal variability in atmospheric gas (a) NH_3 , (c) HNO_3 , (e) SO_2 , (g) HCl and aerosol concentrations (b) pNH_4^+ , (d) pNO_3^- , (f) pSO_4^{2-} , (i) pCl^- , (j) pNa^+ (p in front of component name denotes particulate). Each data point is the monthly averaged concentrations of grouped sites for the period 2006 to 2010, classified according to four ecosystem types: crops ($n = 10$), grassland ($n = 10$), semi-natural ($n = 11$), and forests ($n = 35$). Graph (h) shows the monthly mean ratio of molar-equivalent (equiv.) concentrations of NO_3^- to sum ($\text{NO}_3^- + \text{SO}_4^{2-}$). Month 1: January; Month 12: December.

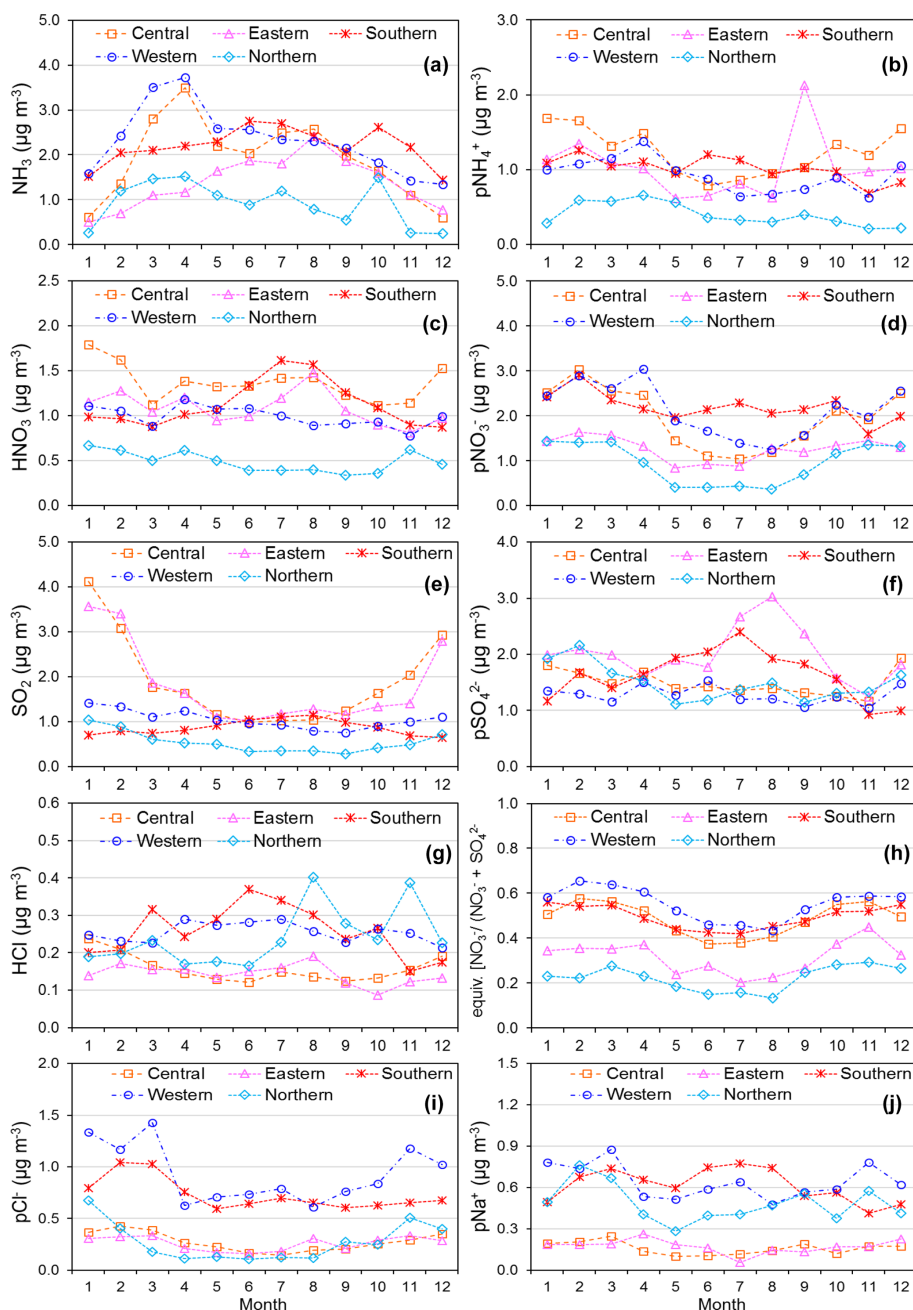


Figure 15. Seasonal variability at sites grouped according to European regions in atmospheric gas (a) NH_3 , (c) HNO_3 , (e) SO_2 , (g) HCl and aerosol concentrations (b) pNH_4^+ , (d) pNO_3^- , (f) pSO_4^{2-} , (i) pCl^- , (j) pNa^+ (p in front of component name denotes particulate). Each data point is the monthly averaged concentrations of grouped sites for the period 2006 to 2010, classified according to five European regions: central ($n = 17$), eastern ($n = 2$), northern ($n = 11$), southern ($n = 12$), and western ($n = 26$). Graph (h) shows the monthly mean ratio of molar-equivalent (equiv.) concentrations of NO_3^- to sum ($\text{NO}_3^- + \text{SO}_4^{2-}$). Month 1: January; Month 12: December.

forest sites, reflecting the elevated regional background in NH_3 concentrations located within agricultural landscapes. This rises rapidly with improving weather conditions and peaks in the spring to coincide with the main period for manure spreading and fertilizer application before the sowing of arable crops (Hellsten et al., 2007). The distinct spring-

time maxima in NH_3 also reflects implementation of the Nitrates Directive (EC, 1991), which prohibits manure spreading in winter. In summer, the second peak in NH_3 concentrations may be associated with increased land surface emissions promoted by warm, dry conditions and possibly with the application of fertilizers. The smaller autumn peak is also

expected to be related to seasonal farming activities (e.g. manure spreading). The key drivers for seasonal variability in NH₃ concentrations at crops sites are therefore a combination of seasonal changes in agricultural practices (e.g. timing of fertilizer/manure applications) and climate that will affect emissions, concentrations, transport, and deposition of NH₃.

Grassland sites

An additional major source of NH₃ in this group of sites is expected to come from grazing emissions and housed livestock (e.g. cattle). Concentrations in this group of sites were generally 2–3 times larger than semi-natural sites (Fig. 14a), attributed to the increased emissions and concentrations from livestock (Hellsten et al., 2007). The spring peak is related to the practice of fertilizer and manure being spread on grazing fields to aid spring grass growth, which will be cut for hay and silage later in the year. NH₃ concentrations in June and July are smaller than in spring or late summer, possibly because grass will be actively growing with possible uptake and removal of NH₃ from the atmosphere (Sutton et al., 2009). The concentrations are also larger in summer than winter, with warmer conditions promoting NH₃ volatilization and thermodynamic shift in NH₄NO₃ to the gas phase.

European regions

The seasonal profiles of NH₃ for central and western European regions were similar, characterized by a large peak in spring that is likely to be agriculture-related (Fig. 15a), as observed at cropland sites (Fig. 14a). While the peak concentrations in both regions are of comparable magnitude (central = 2.6 µg NH₃ m⁻³, western = 2.8 µg NH₃ m⁻³), winter concentrations in central Europe (0.6 µg NH₃ m⁻³) were 3 times smaller than the west (1.5 µg NH₃ m⁻³). This may be related to lower regional background in NH₃ concentrations and/or suppressed emissions in colder temperatures of central Europe in winter. By contrast, eastern and southern European regions have a broad peak in summer, although the eastern region also has a second peak in October (likely agriculture-related). The smallest concentrations were found in northern Europe, with the lowest NH₃ emissions (Fig. 9). The three peaks in the profile show elevated concentrations in summer driven by warming temperatures, with the spring and autumn peaks attributed to influence from NH₃ emissions from agricultural sources.

3.5.2 HNO₃

The seasonal distribution in HNO₃ is similar between the different ecosystem groups, varying only in magnitude of concentrations (Fig. 14c), and reflects the secondary nature of this component that is formed from oxidation of NO_x (Fahney et al., 1986; ROTAP, 2012). Since the HNO₃ data are actually the sum of HNO₃ and HONO, with a small contribution from NO₂ (see Sect. 2.2.3), the temporal patterns seen

are likely to be the superimposed profiles of both HNO₃ and HONO. In the studied region, NO₂ is predominantly from vehicular emissions, which are not expected to show large seasonal variations and should therefore exert a negligible effect on the temporal patterns in HNO₃. With this caveat in mind, HNO₃ concentrations in the crops group are up to 2 times larger than the grassland group, while the smallest concentrations are in the semi-natural group. This is likely related to the proximity of sites in the different groups to combustion sources. A weak seasonal cycle is seen in the secondary HNO₃ air pollutant in all cases, with slightly higher concentrations in late winter, spring, and summer and lowest in March and November. The reaction of NO₂ with the OH radical is an important source of HNO₃ during daytime, whereas N₂O₅ hydrolysis is considered an important source of HNO₃ at nighttime (Chang et al., 2011). Larger HNO₃ concentrations in summer are therefore from increased OH radicals for reaction with NO₂ to form HNO₃. Similarly, higher concentrations of ozone in spring in Europe (EMEP, 2016) can potentially increase HNO₃ concentrations in springtime. Conversely, HNO₃ concentrations are lower in winter, when oxidative capacity is less.

Seasonal variability in HNO₃ will also be influenced by gas–aerosol-phase equilibrium. In the atmosphere, HNO₃ reacts reversibly with NH₃, forming the semi-volatile NH₄NO₃ aerosol if the necessary concentration product [HNO₃] · [NH₃] is exceeded (Baek et al., 2004; Jones and Harrison et al., 2011). Because of this process, the prime influences upon HNO₃ concentrations at sites where NH₄NO₃ is formed are expected to be ambient temperature, relative humidity, and NH₃ concentrations that affect the partitioning between the gas and aerosol phase (Allen et al., 1989; Stelson and Seinfeld, 1982). The availability of surplus NH₃ in spring (Sect. 3.5.1) would tend to reduce HNO₃ and increase NH₄NO₃ formation, which is reflected in the reduced HNO₃ concentrations observed in March, when NH₃ is at a maximum. In summer, warmer, drier conditions promote volatilization of the NH₄NO₃ aerosol, increasing the gas-phase concentrations of HNO₃ and NH₃ relative to the aerosol phase. Seasonality in HNO₃ is therefore complex, related to traffic and industrial emissions, photochemistry, and HNO₃ : NH₄NO₃ partitioning.

An analysis of the same data grouped according to geographical region revealed distinctive cycles in HNO₃ in eastern and southern Europe (Fig. 15c). These two regions showed the highest concentrations in summer and lowest in winter, consistent with enhanced photochemistry in warmer, sunnier climates, and thermodynamic equilibrium favouring gas-phase HNO₃ (Fig. 15c). Summertime peak concentrations in NH₃ were also observed in these two regions (Fig. 15a). In comparison, the seasonal profiles of HNO₃ in other regions were similar to those described for different ecosystem types (Fig. 14c).

3.5.3 SO₂

Seasonality in SO₂ shows concentrations peaking in winter at most sites (Fig. 14e), except in southern Europe, where the peak appeared in summer (Fig. 15e). Increased SO₂ emissions from combustion processes (heating) in the winter months, coupled to stable atmospheric conditions, can result in build-up of concentrations at ground level, thereby contributing to the peak wintertime concentrations. The largest winter concentrations in central and eastern regions exceeded summer values on average by a factor of 4 compared with smaller differences in other regions (Fig. 15e). Enhanced oxidation processes in summer also tend to further reduce concentrations of SO₂ through the oxidation of SO₂ to H₂SO₄ (Saxena and Seigneur, 1987; Sickles and Shadwick, 2007; Paulot et al., 2017). In southern Europe, the seasonal cycle has winter minima and summer maxima instead, likely from increased combustion sources to meet energy demands for air conditioning over the hot summer months. Section 3.4 shows that SO₂ was spatially correlated to HNO₃; differences in relative concentrations between the different ecosystem groups (Fig. 14e) are thus also likely related to relative distance from emission sources.

3.5.4 NH₄⁺, NO₃⁻, and SO₄²⁻

The seasonal profiles of particulate NH₄⁺ (Figs. 14b and 15b) were mirrored by particulate NO₃⁻ (Figs. 14d and 15d) in all groups, demonstrating temporal as well as regional (see Sect. 3.3.4) correlation between these two components. Since NH₄NO₃ is more abundant than (NH₄)₂SO₄, the seasonality of NH₄⁺ is likely to be influenced more by the temperature and humidity dependence of the semi-volatile NH₄NO₃ than by the stable (NH₄)₂SO₄. In summer, warmer and drier conditions promote the dissociation of NH₄NO₃, decreasing particulate-phase NH₄NO₃ relative to gas-phase NH₃ and HNO₃. This process accounts for the summertime minima in NH₄⁺ (Figs. 14b and 15b) and NO₃⁻ (Figs. 14d and 15d). Conversely, cooler temperatures and higher-humidity conditions in winter, spring, and autumn shift the equilibrium to the aerosol phase, with observed peaks in concentrations of NH₄⁺ and NO₃⁻. Since NH₃ concentrations are also generally higher in spring than in autumn (Figs. 14a, 15a), the increased availability of NH₃ in this period contributes towards the higher concentrations of NH₄NO₃ in spring than in autumn. In winter, the combination of NH₄NO₃ remaining in the aerosol phase with the stable conditions that can often develop maintains high concentrations of NH₄⁺ and NO₃⁻ in the atmosphere. The peak in NO₃⁻ in southern Europe was in February only, compared with broader peaks (February–April) in other regions (Fig. 15d), which may reflect differences in climatic conditions. In Figs. 14h and 15h, the ratios of the molar-equivalent concentrations of NO₃⁻ to sum (NO₃⁻ + SO₄²⁻) are plotted. The ratios were highest in spring

and autumn and lowest in summer, lending support to the importance of NH₄NO₃ in controlling the seasonality of NH₄⁺.

In the seasonal profiles for particulate SO₄²⁻, clear summer maxima and winter minima were observed at sites in southern and eastern Europe (Fig. 15f). The peaks occurred at different times, in July (southern Europe) and in August (eastern Europe) (Fig. 15f), and coincided with the timing of corresponding peaks in NH₃ concentrations (Fig. 15a), illustrating the importance of NH₃ in driving the formation of the stable (NH₄)₂SO₄. Since (NH₄)₂SO₄ is formed through the preferential and irreversible reaction between the precursor gases (Bower et al., 1997), particulate SO₄²⁻ concentrations will be governed by the availability of NH₃ and H₂SO₄ (from oxidation of SO₂). As discussed earlier, SO₂ concentrations in southern Europe have a different seasonal cycle from other regions, with higher concentrations in summer than in the winter months (Fig. 15e). Although the seasonal cycle for eastern Europe showed the smallest SO₂ concentrations in the summer, the summer minima here (mean = 1.3 µg SO₂ m⁻³) are in fact larger than the summer peak in southern Europe (mean = 1.1 µg SO₂ m⁻³) and concentrations in other regions (0.4–1.0 µg SO₂ m⁻³). Enhanced summertime concentrations in HNO₃ were observed in these two regions (Fig. 15b), which also suggests potentially increased oxidative capacity for more of the SO₂ to be converted to H₂SO₄ (Sect. 3.5.3). The ready availability of both SO₂ (and conversion to H₂SO₄) and NH₃ (Fig. 15a) in southern and eastern regions in this period thus coincides to produce the summer peak in particulate SO₄²⁻.

In other regions (central, northern, western), formation of (NH₄)₂SO₄ will be limited by the availability of SO₂, which is lowest in summer (Fig. 15e). Conversely, SO₂ concentrations are highest in winter (Fig. 15e), but lower oxidative capacity at this time of year limits formation of H₂SO₄. Since NH₃ concentrations are also smallest in winter (Fig. 15a), formation of (NH₄)₂SO₄ is also limited in winter. This accounts for the higher concentrations of particulate SO₄²⁻ concentrations in winter and in early spring in these regions (Fig. 15f).

3.5.5 HCl, Cl⁻, and Na⁺

The concentrations of HCl measured at all sites and in all groups were very small, with monthly mean concentrations varying between 0.1 and 0.3 µg HCl m⁻³ (Figs. 14g and 15g). There is no discernible seasonality in the data, which suggests that either sites in the network are not affected by any large sources of HCl or that small differences between months are not detectable due to measurement uncertainties at the very low concentrations (method limit of detection ~ 0.1 µg HCl m⁻³ for monthly sampling). By contrast, Cl⁻ (Figs. 14i and 15i) has a distinctive seasonal cycle with higher concentrations in the winter months than summer, similar to that of Na⁺ (Figs. 14j and 15j). The temporal correlation in the data therefore lends further support that Na⁺

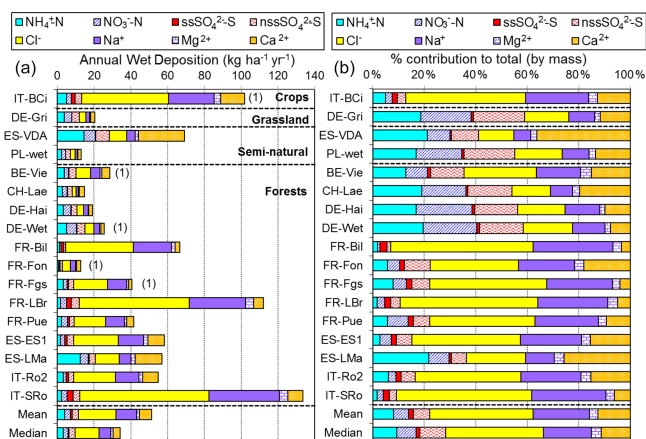


Figure 16. (a) Annual wet deposition of inorganic components ($\text{kg ha}^{-1} \text{yr}^{-1}$) estimated from Rotenkamp bulk precipitation collectors in the NEU bulk wet-deposition network. (b) Percentage contribution of inorganic components to total (by mass) measured at 17 sites from 2008 to 2010. The data shown are 2-year-averaged deposition, made between 2008 and 2010, except at five sites with 1 year of measurement only, as indicated in the graph in brackets.

and Cl^- in the measurements are mainly sea salt (see also spatial correlation in Sect. 3.4). The highest concentrations of Na^+ and Cl^- during winter months would be consistent with increased generation and transport of sea salt generated by more stormy weather from marine sources during those periods (O'Dowd and de Leeuw, 2007).

3.6 Bulk wet-deposition measurements

Annual mean wet deposition of chemical species measured at the NEU bulk sampling sites was estimated by combining measured concentrations with annual precipitation. Site changes also occurred during the operation of the bulk wet-deposition network, with some sites closed and new sites added. At Mitra (PT-Mi3), contamination of the rain samples from bird strikes resulted in the rejection of a large proportion of the monthly data, and this site was excluded from the data analysis. In total, 12 sites provided 2 years of monthly data, with a further 5 sites providing 1 year of monthly data over the period 2008 to 2010. Due to differences in start and end dates for bulk measurements between the sites, the annual mean data derived are for 12-month periods or 2×12 -month periods and not from calendar years.

Annual mean wet-deposition data for the 17 sites from seven countries (Belgium, France, Germany, Italy, Poland, Spain, and Switzerland) are summarized in Fig. 16. Using Na^+ as a tracer for sea salt (Keene et al., 1986), nss-SO_4^{2-} concentrations were also estimated from the total SO_4^{2-} (see Sect. 2.2.2) and are included for comparison. Since the measurements were made at a limited number of sites across Europe, there is insufficient information to make inferences about spatial differences in concentrations. Detailed assess-

ments of extensive precipitation chemistry across Europe are made elsewhere, for example from the EMEP wet-deposition networks (EMEP, 2016; Trseth et al., 2012). What the NEU bulk network data clearly shows is that N_r components in rain also exceed those of S (Fig. 16), as was observed in the atmospheric data. The mean proportional contribution of total N (NH_4^+ and NO_3^-) to the sum total of all wet-deposited species measured (by mass) was 19 % (range = 3 %–39 %), compared with a smaller 9 % (range = 1 %–19 %) contribution from nss-SO_4^{2-} (Table S14). Wet-deposited N (NH_4^+ and NO_3^-) was on average 2 times higher than nss-SO_4^{2-} , similar to that seen in the relative proportion of total N_r (sum of NH_3 , NH_4^+ , HNO_3 , NO_3^-) to total S (sum of SO_2 , SO_4^{2-}) in the atmospheric data (Sect. 3.3.4). Similar to the atmospheric data (Sect. 3.3.4), a considerable fraction of the wet-deposited components was made up of sea salt (Na^+ and Cl^-), with the sum of Na^+ and Cl^- contributing on average 50 % of the total wet-deposited components (range = 20 %–84 %, $n = 17$). Contributions by the other base cations Ca^{2+} and Mg^{2+} gave a further 20 % (range = 8 %–41 %, $n = 17$) (Table S14).

The wet-deposition data on NH_4^+ and NO_3^- , combined with a wider precipitation chemistry dataset (e.g. from EMEP and other national precipitation networks), were used to estimate total N_r deposition to a site (Flechard et al., 2011, 2020). Together, the dry (DELTA[®] network) and wet N_r estimates (NEU bulk network, combined with data from other national precipitation chemistry networks) are used to compare with EMEP models and to examine the interactions between N_r supply and greenhouse gas exchange at the NEU DELTA[®] sites, presented in a separate paper by Flechard et al. (2020). The wet-deposition measurements in this paper highlight where DELTA[®] and bulk wet-deposition data are co-located and provide parallel information on gas and aerosol concentrations (for dry-deposition modelling) and wet deposition at those sites. The co-located data are important for deriving N budgets and linking to ecosystem response (e.g. Flechard et al., 2020) and invaluable for modellers.

4 Conclusions

The NitroEurope DELTA[®] network has provided for the first time a comprehensive quality-assured multi-annual dataset on reactive gases (NH_3 , HNO_3 , SO_2 , HCl) and aerosols (NH_4^+ , NO_3^- , SO_4^{2-} , Cl^-) across the major gradients of emission densities, ecosystem type, and climatic zones of Europe. By sharing the method and protocol with several European laboratories and developing synergies with established infrastructure (e.g. CarboEurope network and EMEP field sites), it has proven possible to establish a large-scale network within a relatively short timescale and with low costs. Key elements were a harmonized methodology and the implementation of quality protocols that included regular lab-

oratory and field inter-comparisons to monitor and improve performance.

At the same time, the concurrent measurement of the gas and aerosol components permitted an assessment of the atmospheric composition and the spatial and seasonal characteristics in the gas and aerosol phase of these components. The dataset has also been used to develop estimates of site-based N_r dry-deposition fluxes across Europe, including supporting the development and validation of long-range transport models. Combined with estimates of wet deposition (from NEU bulk wet-deposition network and other networks such as EMEP), an assessment of the interactions between N supply and greenhouse gas exchange was addressed in a separate paper by Flechard et al. (2020) using N_r and CO₂ flux data from the co-location of the NEU DELTA[®] with CarboEurope Integrated Project sites.

Two key features have emerged in the data. The first is the dominance of NH₃ as the largest single component at the majority of sites, with molar concentrations exceeding those of HNO₃ and SO₂ combined. As expected, the largest NH₃ concentrations were measured at cropland sites, in intensively managed agricultural areas dominated by NH₃ emissions. The smallest concentrations were at remote semi-natural and forest sites, although concentrations in the Netherlands, Italy, and Germany were up to 45 times larger than similarly classed sites in Finland, Norway, and Sweden ($< 0.6 \mu\text{g NH}_3\text{-N m}^{-3}$), illustrating the high NH₃ concentrations that sensitive habitats are exposed to in intensive agricultural landscapes in Europe. The second key feature is the dominance of NH₄NO₃ over (NH₄)₂SO₄, with on average twice as much NO₃⁻ as SO₄²⁻ (on a molar basis). A change to an atmosphere that is more abundant in NH₄NO₃ will likely increase the atmospheric lifetimes and extend the footprint of the NH₃ and HNO₃ gases by the re-volatilization of NH₄NO₃ in warm weather.

Temporally, peak concentrations in NH₃ for crops and grassland sites occurred in spring, reflecting the implementation of the EU Nitrates Directive that prohibits winter manure spreading. The spring agriculture-related peak was seen even at semi-natural and forest sites, highlighting the influence of NH₃ emissions at sites that are more distant from sources. Summer peaks, promoted by increased volatilization of NH₃ but also by gas–aerosol-phase thermodynamics under warmer, drier conditions, were seen in all ecosystem groups, except at forest sites. The seasonality in the NH₃ concentrations thus provided important insights into both the relationship to occurrence of emissions and possible abatement measures to target peak emission periods.

Seasonality in the other gas and aerosol components is also driven by changes in emission sources, chemical interactions, and changes in environmental conditions influencing partitioning between the precursor gases (SO₂, HNO₃, NH₃) and secondary aerosols (SO₄²⁻, NO₃⁻, NH₄⁺).

Seasonal cycles in SO₂ were mainly driven by emissions (combustion), with concentrations peaking in winter, except

in southern Europe, where the peak occurred in summer. HNO₃ concentrations were more complex, as seasonal variations were affected by photochemistry, meteorology, and gas–aerosol-phase equilibrium. Southern and eastern European regions provided the clearest seasonal cycle for HNO₃, with the highest concentrations in summer and the lowest in winter, attributed to increased photochemistry in the summer months in hotter climates. In comparison, a weaker seasonal cycle is seen in other regions, with marginally elevated concentrations in late winter, spring, and summer and lowest in March and November. Increased ozone in spring is likely to enhance oxidation of NO_x to HNO₃ for forming the semi-volatile NH₄NO₃ by reaction with a surplus of NH₃. Cooler, wetter conditions in spring also favour the formation of NH₄NO₃, and more of the NH₄NO₃ remains in the aerosol or condensed phase. This accounts for the higher concentrations of NH₄⁺ and NO₃⁻ in spring and the absence of a HNO₃ peak at this time of year. Conversely, increased partitioning to the gas phase in summer decreases NH₄NO₃ concentrations relative to gas-phase NH₃ and HNO₃. Particulate SO₄²⁻ showed large peaks in concentrations in summer in southern and also eastern Europe, contrasting with much smaller peaks occurring in early spring in other regions. The peaks in particulate SO₄²⁻ coincided with peaks in NH₃ concentrations, illustrating the importance of NH₃ in driving the formation of (NH₄)₂SO₄. Since NH₄NO₃ is more abundant than (NH₄)₂SO₄, the seasonality of NH₄⁺ is likely to be influenced more by the temperature and humidity dependence of the semi-volatile NH₄NO₃ than by the stable (NH₄)₂SO₄. This is supported by similarity in the seasonal profiles of NH₄⁺ and NO₃⁻ at all sites, demonstrating temporal as well as regional correlation between these two components.

Data from the network showed that critical levels of 1 and 3 $\mu\text{g NH}_3 \text{ m}^{-3}$ for the protection of lichens and bryophytes and vegetation were exceeded at 62 % and 27 % of the sites, respectively. At the same time, NH₃ dry deposition will also contribute to a significant fraction of deposited acidity and total N deposition to sensitive habitats, along with NH₄⁺ and HNO₃ dry deposition and wet-deposited NH₄⁺ and NO₃⁻. Although the concentrations of SO₂ have fallen to very low levels at all sites ($< 1 \mu\text{g SO}_2\text{-S m}^{-3}$), SO₂ will continue to be important in contributing to the exceedance of acidification in European ecosystems (EEA, 2019) since SO₂ has a higher acidification potential than NO_x (0.70 kg SO₂ = 1 kg equivalent NO₂ in acidity) (see Hauschild and Wenzel, 1998). Changes in the relative concentrations of the pollutant gases captured in the data suggest that the deposition rates of SO₂ and NH₃ will increasingly be controlled by the molar ratio of NH₃ to combined acidity (sum of SO₂, HNO₃, and HCl), and deposition models should take these changes into account. Indications from the current and projected trends in emissions of SO₂, NO_x, and NH₃ are that NH₃ and NH₄NO₃ will continue to dominate the inorganic pollution load over the next decades, contributing to ecosystem effects through acid and N deposition. The growing relative importance of

NH₃ and NH₄⁺ to total acidic and total N deposition indicates that strategies to tackle acidification and eutrophication need to include measures to abate emissions of NH₃ (Sutton and Howard, 2018).

There is still a lack of NH₃ and speciated monitoring of the inorganic gas and aerosol composition across the EU. An implementation of the DELTA[®] approach across Europe would provide cost-efficient monitoring of the gas- and aerosol-phase pollutants for which reduction commitments are set out in Annex II to the NECD. Monitoring of NH₃ and the interacting acid gases and aerosols is needed to assess contributions of NH₃ to PM_{2.5}, known to be harmful to human health. In addition, such monitoring will also provide the baseline and evidence against which any changes and potential recovery in ecosystem response to changes in emissions of the pollutant gases can be assessed, as required under Article 9 of the NECD. Issues such as human health impacts from fine ammonium aerosols will also drive policy decisions since controlling NH₃ should also reduce PM concentrations (Backes et al., 2016).

Data availability. Summary data are provided in the Supplement. The full dataset is available from <http://www.nitroeuropa.ceh.ac.uk/> (Tang et al., 2020).

Supplement. The supplement related to this article is available online at: <https://doi.org/10.5194/acp-21-875-2021-supplement>.

Author contributions. YST coordinated the establishment of the networks, measurement, and collection of data with the support of several European laboratories. A large number of research institutes provided monitoring sites and local support for installation of equipment and carrying out the monthly exchange of samples. MAS conceived the NEU project and the DELTA[®] network. IS helped with designing and building the low-voltage DELTA[®] equipment. EN helped with network logistics and provided science advice. NvD helped with running proficiency testing schemes and inter-comparisons. UlricD provided GIS support and science advice. CP, MJS, and UlricD facilitated and hosted the DELTA[®] inter-comparisons at their field sites. UlricD also sourced Rotenkamp bulk collectors for the project. JNC provided advice on bulk wet-deposition measurements and calculations. CRF, UlricD, SV, MM, HTU, and MJS led the chemical laboratories that shared the DELTA[®] and wet-deposition measurements. DL developed the NEU database and provided support in data submission. Several of the authors contributed to measurements, network operations, and equipment and site maintenance. YST performed all the data collection and data analysis (including statistics) and wrote the manuscript, with input from all co-authors. MAS, MRH, CRF, UlricD, MF, and JNC provided valuable advice on the interpretation of results and feedback on the manuscript.

Competing interests. The authors declare that they have no conflict of interest.

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